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Time intervals and distances travelled for prehospital ambulance stroke care: data from the randomised-controlled ambulance-based Rapid Intervention with Glyceryl trinitrate in Hypertensive stroke Trial-2 (RIGHT-2)

Mark Dixon,1,2 Jason P Appleton,3 Polly Scutt,1 Lisa J Woodhouse,1,4 Lee J Haywood,1 Diane Havard,1 Julia Williams,1,4 Aloysius Niroshan Siriwardena,1,5 Philip M Bath,1 on behalf of the RIGHT-2 Investigators

ABSTRACT

Objectives Ambulances offer the first opportunity to evaluate hyperacute stroke treatments. In this study, we investigated the conduct of a hyperacute stroke study in the ambulance-based setting with a particular focus on timings and logistics of trial delivery.

Design Multicentre prospective, single-blind, parallel group randomised controlled trial.

Setting Eight National Health Service ambulance services in England and Wales; 54 acute stroke centres.

Participants Paramedics enrolled 1149 patients assessed as likely to have a stroke, with Face, Arm, Speech and Time score (2 or 3), within 4 hours of symptom onset and systolic blood pressure >120 mm Hg.

Interventions Paramedics administered randomly assigned active transdermal glyceryl trinitrate or sham. Ambulance-based stroke trials in urban and rural locations.

Trial registration number ISRCTN26986053.

STRENGTHS AND LIMITATIONS OF THIS STUDY

⇒ The first multicentre paramedic-delivered ambulance-based randomised controlled trial in stroke in the UK.
⇒ Ambulance response time intervals and distances are collated and reported for 1149 patients assessed as likely to have a stroke.
⇒ The time interval between arrival at hospital and the ambulance becoming available for the next emergency call (hospital turnaround) is not captured, but worth considering for future trials.
⇒ Timing and logistic data may not be fully representative of all urban and rural locations due to non-participation of some hospitals and ambulance stations within ambulance service regional areas.

INTRODUCTION

Routine prehospital management of suspected acute stroke involves rapid identification of suspected stroke using a validated stroke screening tool, prompt transport, pre-arrival notification and primary stabilisation to the nearest appropriate receiving stroke centre.1 The mainstays for hyperacute management of stroke in hospital include urgent neuroimaging, stroke unit care, reperfusion therapy for ischaemic stroke and blood pressure (BP) lowering for intracerebral haemorrhage.2 For reperfusion therapies, shortening the time from symptom onset to treatment improves functional outcome and this has become the

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For numbered affiliations see end of article.

Correspondence to Philip M Bath; philip.bath@nottingham.ac.uk

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aim of prehospital and in-hospital acute stroke services.3–6
Thus, ambulance services play a crucial role in assessing, identifying and conveying patients with suspected stroke to primary and comprehensive stroke centres, which may include bypassing local emergency departments.

Timely prehospital care for stroke is dependent on several factors that include rapid recognition of potential stroke and calling for help,7 ambulance response times encompassing symptom onset to arrival at hospital,8 distance from scene to hospital9 and the accuracy of identifying patients with true stroke or transient ischaemic attack from those with a stroke mimic.10 There are a small, but growing number of studies that explore randomised paramedic-initiated interventions commencing in the ambulance for acute stroke. However, few studies have systematically analysed these parameters and the factors that influence them in acute prehospital stroke practice.11–12

Here, we report the logistics underlying patient recruitment to the Rapid Intervention with Glyceril trinitrate in Hypertensive stroke Trial-2 (RIGHT-2), a large ambulance-based stroke trial in the UK that investigated the efficacy of transdermal glyceril trinitrate as a paramedic-delivered intervention in suspected acute stroke. Specifically, ambulance response times and distance travelled across multiple organisations in this setting are assessed.

METHODS
RIGHT-2 trial
RIGHT-2 commenced recruitment in September 2015 with the first participant recruited on 22 October 2015.

RIGHT-2 was a multicentre prospective, single-blind, parallel group randomised controlled trial; the protocol, statistical analysis plan, baseline data, main results and subgroup results in participants with a final diagnosis of intracerebral haemorrhage are published.13–17 Briefly, adult patients with suspected stroke presenting to the emergency service via an emergency call were recruited if they were FAST-positive (facial weakness, arm weakness, speech abnormality; with test score 2 or 3), had systolic BP of >120 mm Hg, were within 4 hours of symptom onset, presented to trial-trained paramedics from eight UK ambulance services and were to be taken to a trial-participating hospital. Patients were randomised to receive transdermal glyceril trinitrate (GTN) or sham patch in the ambulance and this was continued for three further days during hospital admission.13 The study was undertaken across eight UK ambulance services (AS): East of England AS (EEAS), East Midlands AS (EMAS), London AS (LAS), South-Central AS (SCAS), South-West AS (SWAS), Welsh AS (WAS), West-Midlands AS (WMAS) and Yorkshire AS (YAS). All participating ambulance services used FAST identification and protocols consistent with national guidelines.

For each eligible patient, the enrolling paramedic assessed capacity and obtained patient or proxy consent (from a relative on the scene, or from the paramedic witnessed by a colleague), completed a written case report form to capture in-ambulance baseline and on-treatment data and applied the transdermal patch of GTN or sham dressing.13 Ambulance-related data not recorded at source were confirmed by research paramedics from participating ambulance services after review of control room timing logs or patient care records, and then entered into the trial database.

Timings and distances
Timings were obtained from each ambulance service (time of emergency call, resource dispatch, scene arrival and departure, hospital arrival) and from paramedic records (consent for trial enrolment, randomisation, application of study treatment). Paramedic-documented history provided the time of symptom onset or, where unclear, the last known well time.

Distance measurements were calculated from the address or postcode of the emergency location, where available, to the expected stopping point for the ambulance at the destination hospital (accident and emergency or stroke unit entrance) to the nearest 10 metres using Google Maps; one ambulance service was unable to provide postcode information due to time constraints. One ambulance service was able to provide the linear distance from the location of the ambulance at the point of dispatch to the scene of the emergency.

A comparison of urban versus rural ambulance services arbitrarily divided ASs by <25% rural versus >25% rural (as defined in table 1; online supplemental table I).

Comparison of trial and non-trial patients
One ambulance service provided response time interval and distance data for a cohort (n=49) of patients with confirmed stroke who were not enrolled into RIGHT-2 (attended by non-trial trained paramedics) but were transported to the same specialist stroke centres participating in the trial.

Statistical analysis
Time intervals (in min), distances (in km) and baseline characteristics were compared between ambulance services using \( \chi^2 \) and Kruskal-Wallis (one-way analysis of variance on ranks) tests. Multiple comparison procedures (Dunn’s with Bonferroni correction) were used to assess which ambulance service differed from the others. Spearman and point-biserial correlations were performed to identify the relationship between baseline variables, times and distances. Data are number (%), median (IQR) or mean (SD). Statistical significance was defined overall at p<0.05, and at p<0.001 for correlation matrices and multiple comparisons. Statistical analyses were conducted with SPSS V.24 (IBM, New York, USA).

Patient and public involvement
This study was supported by public members of the trial steering committee who were involved throughout, including in trial design, development, conduct, periodic review and dissemination of results.
RESULTS

RIGHT-2 recruited 1149 patients between September 2015 and May 2018. Table 1 outlines patient recruitment across the various participating ambulance services, which collectively covered an area of 122 065 km² in England and Wales (ie, 42% of the land area of these countries). Ambulance services varied considerably in size (1605 km² vs 25 899 km²), population served per service (2.9 million vs 8.6 million) and annual stroke events (7400 vs 13 000) (table 1). Altogether 1492 paramedics volunteered to be trained in the trial, of whom 516 (36%) recruited at least one patient. Where two or more trial paramedics were present at the scene, the paramedic initiating randomisation was credited. On average, 2.2 patients were recruited by each paramedic who enrolled at least one patient although this varied between ambulance services (1.1 vs 3.1).

Patient characteristics

Of the 1149 patients recruited, average age was 73 (15) years, women 48%, BP 162 (25)/92 (18) mm Hg, Glasgow Coma Scale (GCS) 13.9 (1.7) and FAST score of three 60% (online supplemental table II). The final diagnosis varied between ambulance services, with the rate of conditions mimicking acute neurovascular disease ranging from 14.3% to 36.1%. This is consistent with other prehospital trials without physician presence or mobile stroke unit care, and the rate of stroke mimic reported here is explored elsewhere. Baseline temperature also varied. Otherwise, baseline characteristics did not differ between ambulance service. As age increased, BP and glucose were higher, and heart rate, FAST and GCS lower (online supplemental table III). Informed consent was provided by 603 (53%) patients, 431 (38%) relatives and 115 (10%) paramedics witnessed by a colleague on scene.

Time intervals

The time intervals for various stages in the journey from stroke scene to hospital are shown in online supplemental table IV. Overall, the median time from symptom onset to emergency call was 19 (IQR 5–64) min and this did not differ between ambulance services (online supplemental tables IV and V). The median time from emergency call to ambulance dispatch was 3 (1–7) min and varied between ambulance service (1 min vs 5 min). An ambulance resource arrived at the scene within 8 (5–13) min from being dispatched (and 10 (6–16) minutes if only including RIGHT-2 trained paramedics) with this varying between ambulance service (8 min vs 12 min).

The median time from onset of symptoms to randomisation was 71 (45–116) minutes (table 2, figure 1) and this varied between ambulance service (53 min vs 77 min). Significantly, randomisation occurred within 30 and 60 min of symptom onset in 104 (9.1%) and 491 (42.9%) participants, respectively (table 2). Ambulance resources spent a median of 33 (26–46) minutes on scene, though this varied between ambulance services (29 min vs 43 min) (online supplemental table IV). Importantly, time on scene did not differ significantly when comparing RIGHT-2 patients vs non-RIGHT-2 patients 34 (26–44) and...
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Table 2  Timings: symptom onset to randomisation (OTR) (min). Data are N (%), median (25–75 centile); comparison by Kruskal-Wallis test

<table>
<thead>
<tr>
<th>Min</th>
<th>E&amp;W</th>
<th>EEAS</th>
<th>EMAS</th>
<th>LAS</th>
<th>SCAS</th>
<th>SWAS</th>
<th>WAS</th>
<th>WMAS</th>
<th>YAS</th>
<th>p</th>
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<tbody>
<tr>
<td>OTR</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N (%)</td>
<td>1149</td>
<td>178 (15.5)</td>
<td>218 (19.0)</td>
<td>202 (17.6)</td>
<td>7 (0.6)</td>
<td>265 (23.1)</td>
<td>89 (7.7)</td>
<td>37 (3.2)</td>
<td>153 (13.3)</td>
<td>0.001</td>
</tr>
<tr>
<td>Median (25–75 centile)</td>
<td>71 (45–116)</td>
<td>73 (47–120)</td>
<td>59 (35–100)</td>
<td>77 (51–124)</td>
<td>53 (45–65)</td>
<td>75 (49–107)</td>
<td>75 (48–123)</td>
<td>60 (32–115)</td>
<td>70 (45–118)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>N (%)</td>
<td>104 (9.1)</td>
<td>15 (8.4)</td>
<td>38 (17.4)</td>
<td>11 (5.4)</td>
<td>1 (14.3)</td>
<td>16 (6.0)</td>
<td>7 (7.9)</td>
<td>6 (16.2)</td>
<td>10 (6.5)</td>
<td>0.001</td>
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<td>≤30</td>
<td></td>
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<tr>
<td>31–60</td>
<td>387 (33.8)</td>
<td>63 (35.4)</td>
<td>82 (37.6)</td>
<td>61 (30.2)</td>
<td>3 (42.9)</td>
<td>82 (30.9)</td>
<td>28 (31.5)</td>
<td>13 (35.1)</td>
<td>56 (36.6)</td>
<td>0.001</td>
</tr>
<tr>
<td>61–90</td>
<td>258 (22.5)</td>
<td>32 (18.0)</td>
<td>34 (15.6)</td>
<td>51 (25.2)</td>
<td>0 (0.0)</td>
<td>77 (29.1)</td>
<td>19 (21.3)</td>
<td>5 (13.5)</td>
<td>38 (24.8)</td>
<td>0.001</td>
</tr>
<tr>
<td>91–120</td>
<td>136 (15.1)</td>
<td>25 (14.0)</td>
<td>19 (8.7)</td>
<td>25 (12.4)</td>
<td>0 (0.0)</td>
<td>36 (13.6)</td>
<td>13 (14.6)</td>
<td>5 (13.5)</td>
<td>13 (8.5)</td>
<td>0.001</td>
</tr>
<tr>
<td>121–180</td>
<td>173 (15.1)</td>
<td>30 (16.9)</td>
<td>33 (15.1)</td>
<td>28 (13.9)</td>
<td>0 (0.0)</td>
<td>40 (15.1)</td>
<td>16 (18.0)</td>
<td>6 (16.2)</td>
<td>20 (13.1)</td>
<td>0.001</td>
</tr>
<tr>
<td>181–240</td>
<td>76 (6.6)</td>
<td>12 (6.7)</td>
<td>9 (4.1)</td>
<td>20 (9.9)</td>
<td>0 (0.0)</td>
<td>13 (4.9)</td>
<td>5 (5.6)</td>
<td>2 (5.4)</td>
<td>15 (9.8)</td>
<td>0.001</td>
</tr>
<tr>
<td>&gt;240</td>
<td>15 (1.2)</td>
<td>1 (0.5)</td>
<td>3 (1.4)</td>
<td>6 (3.0)</td>
<td>1 (14.3)</td>
<td>1 (0.4)</td>
<td>1 (1.1)</td>
<td>0 (0.0)</td>
<td>2 (0.7)</td>
<td>0.001</td>
</tr>
</tbody>
</table>


Distances

The median distance travelled from the postcode of the suspected stroke scene to the receiving hospital was 10.0 (4.4–18.4) km, with considerable variation between ambulance services (4.1 km vs 19.9 km) (online supplemental table VIII). Time from scene to hospital was moderately positively correlated with distance from scene to hospital (online supplemental figure I:A-G present geographical distribution of randomisation by ambulance service).

Urban versus rural services

When comparing urban and rural ambulance services (online supplemental table I), there was no difference in receipt of the emergency call to dispatching a resource to scene, nor a difference in onset of symptoms to randomisation. The time spent at scene was marginally longer in rural locations and, as anticipated, both conveyance time and distance to the stroke centre was statistically different.

Comparison of trial and non-trial patients

In the ambulance service with times available for patients not enrolled in the trial, on scene to hospital arrival differed among patients enrolled and not enrolled in RIGHT-2, 10 (0.4–64.7) vs 16 (7.6–24.0) min (online supplemental table VIII). The median distance from dispatch location to scene in the ambulance service with this available (EMAS) was 7.3 km (3.5–12.0).

DISCUSSION

In this large national prehospital trial, 516 paramedics from eight ambulance services across England and Wales successfully recruited 1149 participants and transported them to 54 hospitals. Paramedics assessed and diagnosed suspected stroke, consented patients and initiated...
randomised treatment. Key timings were: onset to emergency call 19 min, onset to scene 40 min, onset to randomisation 71 min, time at scene 33 min, randomisation to hospital 24 min and depart scene to hospital arrival 15 min; all but the first two differed between ambulance services. The average distance travelled by one ambulance service from dispatch location to scene was 7.3 km and 10.0 km from scene to hospital for all participating ambulance services.

Prehospital time intervals in acute stroke have been described previously, but rarely in randomised trials. The symptom onset to randomisation time of 71 min in RIGHT-2 is consistent with two previous UK ambulance-based stroke trials (RIGHT was 55 min and Paramedic Initiated Lisinopril For Acute Stroke Treatment (PIL-FAST) was 70 min) although these were small single centre pilot studies undertaken largely in urban settings. The large US Field Administration of Stroke Therapy - Magnesium trial (FAST-MAG) reported a median of 43 min from symptom onset to receipt of study drug. Nevertheless, these times are all longer than UK multicentre ambulance-based trials outside of stroke, notably the AIRWAYS-2 and PARAMEDIC-2 trials in cardiac arrest. In PARAMEDIC-2, the onset of symptoms to initiation of treatment in the intervention group was just 21.5 min. The most important driver of this difference is most likely shorter onset to call times for patients who had cardiac arrest than for stroke, and suspected stroke may require more complex assessment both by call handlers and by paramedics on scene. Additional contributors are that cardiac arrest is allocated the highest dispatch priority, an immediate response and patients receive immediate trial treatment with emergency waiver of consent.

The explanation for differences in timings is probably multifactorial but the degree of urban versus rural population is one likely explanation. This was apparent for time spent at the scene and both time and distance to hospital. As expected, there were no differences for receipt of call to dispatch, arrival of RIGHT-2 trained paramedic at scene nor onset of symptoms to randomisation.

There are several strengths of this study. First, RIGHT-2 involved 8 of 11 ambulance services in England and Wales. Of those not participating, two were unable to join because they were involved in another ambulance-based stroke trial and the other involved hospitals that were concerned about adversely impacting on recruitment to commercial trials. Among 1492 trained paramedics in RIGHT-2 procedures 516 consented and randomised a large number of participants, adhered to the protocol and completed specific data recording. It is noted that there are marked differences in recruit numbers between ambulance services. This, in part, is accounted for due to low recruitment during the initial recruitment phase requiring broadening of ambulance services from 5 to 8 stroke centres from 30 to 54. Furthermore, recruitment hours initially limited to typical working hours for research staff availability were extended to encompass 24/7 recruitment reflective of real-world ambulance care to not limit participation and maximise inclusion.

Secondly, a small number of stroke centres closed recruitment to ambulances once target numbers of participants had been received and before the end of the recruitment phase highlighting the challenging reliance on dual centres when dealing with research in prehospital stroke.

Second, the consent model applied in RIGHT-2 is unlike any other large-scale ambulance-based studies worldwide to date and builds on previous UK based prehospital stroke pilots. Other prehospital trials in stroke have relied on models of either informed consent, deferred consent or consent by doctor (present or remote).

Stroke is complex due to the varying nature of severity of presentations where patients’ ability to consent in an informed manner to participate in a clinical study should not be overlooked preserving patient autonomy in accordance with the Declaration of Helsinki. Notwithstanding the complexities of emergency presentations that could impact on decision-making, mental capacity or short intervention windows and the impact these situations bring to truly informed patient consent, the combined consent approach in RIGHT-2 acknowledges
patient autonomy without precluding participation from those who are unable to voice their opinion or who lack presence of a proxy to consent on their behalf. Mechanisms to safeguard consent were built into the protocol through reconfirmation of consent once in hospital for both the prehospital and in-hospital elements, respectively, and patient and public representatives were fully embedded within protocol development and steering group oversight of the trial.

Third, the protocol required flexibility and adaptation to align with individual operational processes specific to each ambulance service to ensure successful delivery of the trial. Fourth, detailed logistic information on timing and distances travelled were collected. Last, the results highlight the successful delivery of a simple, ambulance-based intervention with 43% of the patients receiving the intervention within 2 hours of symptom onset without compromising time on scene required to complete additional research activity.

There are also several study limitations. First, it is recognised that not every receiving stroke unit within each ambulance service region could participate in RIGHT-2 due to capacity and competing research (this included concurrent commercial and post-arrival trials). Therefore, it must be considered that the timing and logistic data of participating hospitals may not be fully representative of all urban and rural locations. However, the intention was not to assess the differences between urban and rural settings, but to shed light on the conduct and deliverability of a prehospital intervention in stroke where time and distance may impede access to specialist stroke services. Furthermore, stroke unit hours of operation varied across the 54 centres with a small number of sites not accepting patients outside working hours which impacted paramedics’ decisions to randomise. This reduces the reflection of real-world emergency stroke care. The duration of recruitment varied between regions due to complexities in setting up multicentre research.

Additionally, it is acknowledged that the recruitment criteria were broad which resulted in a higher than anticipated proportion of stroke mimics. To mitigate this, mobile stroke unit care is an emerging field where anticipated proportion of stroke mimics. To mitigate this, mobile stroke unit care is an emerging field where the applicability of future large-scale paramedic-delivered ambulance-based stroke trials in urban and rural locations.

Nevertheless, prehospital time intervals and distances from scene-to-hospital varied by ambulance service and this was, at least in part, explained by the type of urban versus rural population. Although our results may not be generalisable to all ambulance service settings, they do inform future developments in ambulance-based stroke care and provide support to the deliverability of future large-scale multicentre pre-hospital paramedic-delivered ambulance-based acute stroke trials.

Author affiliations
1Division of Mental Health and Clinical Neuroscience, University of Nottingham Faculty of Medicine and Health Sciences, Nottingham, UK
2Leicester, Leicestershire & Rutland Division, East Midlands Ambulance Service NHS Trust, Nottingham, UK
3Stroke, University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK
4Division of Paramedic Science, School of Health and Social Work, University of Hertfordshire, Hatfield, UK
5School of Health and Social Care, University of Lincoln, Lincoln, UK

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Contributors PMB, also chief investigator and guarantor, and MD conceived the study; All authors contributed to the planning, design and conduct. LJW was responsible for data curation. PS and LJW supported with statistical analysis. All authors contributed to the reporting, analysis and interpretation of the results. MD and PMB led the writing of the manuscript with critical revision from JPA, PS, LJW, LJH, DH, JW and ANS.

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Patient and public involvement Patients and/or the public were involved in the design, or conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

Patient consent for publication Not applicable.

Ethics approval RIGHT-2 was approved by the UK regulator (Medicines and Healthcare products Regulatory Agency, reference: 03057/0064/001–001; Eudract 2015–000115–40) and national research ethics committee (IRAS: 167115) and was adopted by the National Institute for Health Research Clinical Research Network. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request.

Individual participant data are shared with the Blood pressure in Acute Stroke Collaboration and Virtual International Stroke Trials Archive.

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ORCID iDs Mark Dixon http://orcid.org/0000-0002-4036-3792 Lisa J Woodhouse http://orcid.org/0000-0002-4472-1999 Julia Williams http://orcid.org/0000-0003-0796-5465 Aloysius Niroshan Sirivardena http://orcid.org/0000-0003-2484-8201

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