**INTRODUCTION**

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The deadline for submitting abstracts is **14 September 2020.**

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**PRESENTATION TYPE**

|  |  |
| --- | --- |
| Presentation type | Scientific |

**YOUR DETAILS**

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| Title | **Click to choose** |
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| Title \* (No more than 15 words) |  |
| List of author(s) and institutions(s) and email addresses \* |  |
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|  |  |
| --- | --- |
| Background and aims \* |  |
| Method \* |  |
| Results \* |  |
| Conclusion \* |  |

In addition, please outline the following in no more than 50 words

|  |  |
| --- | --- |
| How will this research improve life after stroke for stroke survivors? \* |  |

**SUBMITTING YOUR PROPOSAL**

Please tick the box here to confirm the following:

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I declare, that to the best of my knowledge, the only direct or indirect interest I have in both profit and not for profit making organisations potentially related to SAFE are those listed below:

At present or within the last 12 months I have been involved in the following activities (please feel free to add lines as appropriate)

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| Employee |  |  |
| Consultant\* |  |  |
| Principal investigator\*\* |  |  |
| Member of steering committee, advisory board or equivalent body |  |  |
| Investigator (not principal) for the development of a product\*\*\* |  |  |
| Speaker in industry sponsored symposia, press conferences or other meetings |  |  |
| I have received lecture fees and travel expenses (paid to my employer) for lectures at symposia |  |  |

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| I am a board member in the following national/international society(ies) other than SAFE |  |
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\* Consultant: expert who charges a fee (personal, institutional or both) for providing advice or services or both

\*\* Principal investigator: investigator responsible for the coordination of investigators at different sites participating in the (multicenter) trial. This includes investigator initiated studies.

\*\*\* Investigator: physician involved in a clinical trial at a specific site. Can be the leader or the member of the clinical trial team. This includes investigator initiated studies.

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