Table 1. Specific strategies and considerations for involving people with lived experience of selfinjury in research.

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Initial Recruitment	 People with lived experience of self-injury are often keen to contribute to research and break down stigma. The approaches below may help when inviting individuals in the research process and during subsequent interactions: Use existing professional groups to invite people with lived experience (e.g. American Association for Suicidology; Canadian Depression Research and Intervention Network; Western Australian Health Translation Network; INVOLVE). Create specific lived experience components of relevant professional organisations (e.g., International Society for the Study of Self-Injury; American Association for Suicidology, International Association for Suicide Prevention). Online self-injury communities are abundant and highly salient for many individuals with lived experience of self-injury. Posts can be made on relevant platforms to invite participation in research. Use of online chat and meeting software (e.g., Skype, Zoom, Webex) may permit follow-up and facilitate further involvement.
Involvement in	People with lived experience of self-injury can participate in all parts of the
Research	 research process. Their involvement may be helpful in the following ways: People with experience of self-injury can use their own experiences to
	guide priority driven research.
	 People with experience of self-injury can use their stories of stigma in the
	healthcare system to inform treatment provision and service delivery.
	People with lived experience of self-injury are in a unique role to inform
	research design through consideration of the impact of research on
	 participants (e.g., self-injury imagery may be triggering). When conducting research, disclosures during interview studies may break
	down stigma and facilitate rapport.
	 Maximising the reach and impact of findings through involving individuals
	with lived experience in conferences and eliciting their viewpoints.
Ethical	Involvement in research may be upsetting or triggering for some people with a
considerations	history of self-injury. The following considerations may help mitigate such risks:
	• In line with ethical guidelines, ensure clear communication regarding
	participant requirements from the outset <i>and</i> during the research process, provide community-based (local) and online resources to all participants, and
	employ mood augmentation or relaxation techniques to ensure participants
	are not upset upon completion of the study.
	People with lived experience can be reimbursed for their time in a number of
	ways. The INVOLVE guidelines in the UK (see below) outline different levels of
	remuneration depending on the level of involvement.
Helpful	Resources to be consulted/adapted for involving people with lived NSSI
Resources	experience in research:
	 INVOVLE: <u>www.invo.org.uk</u> American Association for Suicidology: <u>www.suicidology.org</u>
	 American Association for Successing: <u>www.successing</u> Canadian Depression Research and Intervention Network: www.cdrin.org
	 Recovery-oriented mental health services: www.health.gov.au
	 BC SUPPORT Unit (bcsupportunit.ca)
	Resources to provide to individuals with lived NSSI experience:

•	 Self-injury Outreach & Support: <u>www.sioutreach.org</u>
•	Shedding Light on Self-injury: <u>www.self-injury.org.au</u>
•	Self-injury and Recovery Research and Resources:
	http://www.selfinjury.bctr.cornell.edu