RCloud Tasking Form - Part B: Statement of Requirement (SoR)

Title of Requirement	Rollable Non-thermal Plasma Microbubble Chamber for Degrading CBW Agents				
Requisition No.	As stated in the RCloud Portal				
SoR Version	1				

1.	Statement of Requirements							
1.1	Summary and Background Information							
	Summary . This project aims to develop a simple, easy-to-use method to destroy both chemical and biological warfare agents in austere environments.							
	Background . This project aims to investigate the feasibility of using plasma-activated air microbubbles to disable and eliminate a broad spectrum of CBW agents in a tactical environment. [REDACTED]							
1.2	Requirement							
	Work Package 1 – Feasibility study of plasma - liquid contacting methods Task 1.1.1 [REDACTED] Task 1.1.2 [REDACTED] The contract will run for 4 months, with the option to conduct follow-on work for 6 months							
1.3	Options or follow on work							
	The Authority may at its sole discretion, exercise its right to activate the consecutive and irrevocable controption period detailed in the Schedule of Requirements. It is agreed that the Authority has no obligation to exercise such options							
	Any work carried out by the Contractor on these options without the prior written approval of the Authority Commercial Services Department will be at the Contractor's own risk.							
	A fully costed proposal for the Options is required.							
	Option 1.							
	Reactor Design							
Design and fabricate a rollable plasma reactor								
	o Task 1.2 [REDACTED]							
	o Task 1.3 [REDACTED]							
	o Task 1.4 [REDACTED]							

microbubbles are able to penetrate the air-liquid interface and also into the liquid layer. The output of the feasibility study will be reviewed by Dstl subject matter experts (SMEs) who will decide if the technology

provides sufficient promise to pursue further.

Option 2

- Work Package 2: Quantify the chemical decomposition rate of chemical warfare agent simulants due to plasma treatment
 - Task 2.1 [REDACTED]
 - o Task 2.2 [REDACTED]
 - o Task 2.3 [REDACTED]
 - o Task 2.4 [REDACTED]
 - o Task 2.5 [REDACTED]

Option 3

- Work Package 3: Demonstrate the capability of plasma generated by a rollable plasma reactor to inactivate biological warfare agent using simulants such as robust [REDACTED]
 - o Task 3.1 [REDACTED]
 - o Task 3.2 [REDACTED]
 - o Task 3.3 [REDACTED]
 - o Task 3.4 [REDACTED]
 - Task 3.5 [REDACTED]

The decision to invoke Option 2 and/or 3 will be made once Option 1 is successfully executed: Assembly of rollable plasma reactor is completed and results show that it functions as expected

1.4 Health & Safety, Environmental, Social, Ethical, Regulatory or Legislative aspects of the requirement

The Contractor's work location must have the appropriate licences to handle the **[REDACTED]** options they propose to work with. The contractor must provide evidence of these licences. In addition, the contractor must provide evidence of competence for all involved, to carry out what is outlined in their proposal. The supplier must provide evidence of a robust safety culture and examples of the processes undertaken to ensure H&S of staff. An example risk assessment could be provided to support this.

1.5	Deliverables & Intellectual Property Rights (IPR)						
Ref.	Title	Due by	Format	Expected classification (subject to change)	What information is required in the deliverable		
Base Period - D1	Feasibility Study Review (FSR)	T0 + 4 months	Written report on customer template	Official	Report on findings including recommendation on feasibility of non-thermal plasma in liquid		
Base Period – D2	Monthly-project progress review	T0 + 1-4 months	MS Teams teleconference or site visit	Official	Review to include but not limited to: • Update on technical progress • Risks/issues		
Option Period – D3	Monthly-project progress review	T0 + 5-10 months	MS Teams teleconference or site visit	Official	Review to include but not limited to: • Update on technical progress • Risks/issues		
Option Period – D4	Report on plasma reactor design	T0 + 6 months	Written report on customer template	Official	Report on findings		
Option Period – D5	Report on chemical simulant testing	T0 + 10 months	Written report on customer template	Official	Report on test results		

Option Period – D6	Report on biological simulant testing	T0 + 10 months	Written report on customer template	Official	Report on test results
Option Period – D7	Practical demonstration	T0 + 10 months	Physical demonstration at contractor site	Official	

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1.7 Deliverable Acceptance Criteria

The report must cover all the areas outlined in 'requirements' in section 1.4 above, and also meet the following requirements:

- The review will be written in a user defined template, acceptable to the authority, to a standard of a peer-reviewed article e.g. publication of a peer reviewed technical journal, or as detailed in the Defence Research Reports Specification (DRRS) which defines the requirements for the presentation, format and production of scientific and technical reports prepared for MoD (https://www.gov.uk/guidance/submit-a-report-to-athena#defence-research-report-specification)
- All reports shall be free from spelling and grammatical errors, containing a full bibliography
 of all the sources used.