# PERFORMANCE DATA ON COLD TEMPERATURE DISPERSION OF CF3I

by

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# **INTRODUCTION**

Trifluoroiodomethane (CF<sub>3</sub>I) has been proposed as a potential replacement for halon 1301 in aircraft engine nacelle and dry bay fire protection applications [Grosshandler *et al.*, 1994; Gann, 1995]. The potential use of CF<sub>3</sub>I in fuel tank ullage inerting has also been considered recently [Tucker, 1998]. Before CF<sub>3</sub>I can be considered as a potential drop-in replacement, several operational issues need to be addressed or re-examined. The proposed work is intended to examine one important aspect related to the application of CF<sub>3</sub>I: cold temperature discharge of CF<sub>3</sub>I into a sub-zero environment.

Table 1 lists some of the physical properties of  $CF_3I$  and  $CF_3Br$ . Since  $CF_3I$  has a normal boiling point of -22 °C, the dispersion of  $CF_3I$  into air at temperatures down to -40 °C may not be as effective as halon 1301 which has a normal boiling point of -57.8 °C. Although discharge of cold  $CF_3I$  (chilled to about -40 °C) into a compartment at ambient room temperature has been examined at NIST [Gann, 1995], the discharge of cold  $CF_3I$  into a cold compartment has not been performed, or at least has not been documented in open literature. In order to assure that there is no substantial deterioration in dispersion performance of  $CF_3I$  under cold temperature applications, discharge tests in a simulated fire compartment at the lowest temperatures expected in service should be conducted.

Table 1. Selected physical properties of CF<sub>3</sub>I and CF<sub>3</sub>Br [from Gann, 1995].

Agent	Molecular weight	$T_b$	$T_c$	$P_c$	$\rho_c$	$\Delta H_{v}$
	(kg/mol)	(°C)	(°C)	(MPa)	$(kg/m^3)$	(kJ/kg)
CF <sub>3</sub> I	0.196	-22.0	122.0	4.04	871	106
CF <sub>3</sub> Br	0.149	-57.8	67.0	4.02	745	111

 $T_b$  is the normal boiling point;  $T_c$  is the critical temperature;  $P_c$  is the critical pressure;  $\rho_c$  is the critical density;  $\Delta H_v$  is the latent heat of vaporization at  $T_b$ .

An example is used to illustrate the importance of studying the discharge and dispersion of  $CF_3I$  in a sub-zero environment. Consider a 3 L container with 4.5 kg of  $CF_3I$  pressurized with nitrogen to 4.12 MPa (600 psia) at 22 °C. If this container with agent is cooled down to -40 °C, the final pressure of the container is estimated to be 2.8 MPa using the computer code (PROFISSY) developed by the National Institute of Standards and Technology [Gann, 1995]. Assuming the discharge of the liquid agent from the container is an isentropic process from the initial container pressure to atmospheric pressure, the final states of the agent can

be calculated using the computer code (FISSYCS), which is a derivative of PROFISSY [Gann, 1995]. Table 2 tabulates the calculated results. Comparing the two initial conditions, the liquid fraction (the percent of agent/nitrogen mixture still remained in liquid phase after the isentropic expansion process) is substantially higher at -40 °C. Such a high liquid fraction should have a significant effect on the subsequent evaporation and dispersion of the agent/nitrogen mixture.

Irrespective of the break-up mechanisms, the liquid core from the discharge nozzle will be disintegrated subsequently into small droplets. At 22 °C, the liquid droplets rapidly evaporate into vapor, thus facilitating the agent dispersion. However, at – 40 °C, the liquid droplets evaporate slowly, and a two-phase (droplet-gas) flow persists longer, thus hindering the effective dispersion of agent droplets especially when baffles are present. This point is further illustrated in Figure 1, which shows the calculated evaporation time as a function of ambient temperature for three initial droplet diameters when the ambient agent mass fraction is zero. The evaporation time was calculated using the classical *d*<sup>2</sup>-law for droplet evaporation [*e.g.*, Chin and Lefebvre, 1983.]. For simplicity and illustrative purposes, neat CF<sub>3</sub>I droplets were used in the calculations. When the ambient agent mass fraction is not zero, the evaporation time of an agent droplet is longer because of the reduction in the concentration gradient for mass transfer. Figure 2 shows the effect of ambient CF<sub>3</sub>I mass fraction on the evaporation time of a 100 μm droplet. A mass fraction of 0.316 corresponds to twice the cup-burner value (mole fraction of 0.032 [Grosshandler *et al.*, 1993]) of CF<sub>3</sub>I. Note that in an actual discharge, the agent droplets will not be evaporating at zero ambient agent mass fraction.

Table 2. Liquid fraction of CF<sub>3</sub>I/N<sub>2</sub> mixture after isentropic expansion to 0.101 MPa.

Initial conditions	Liquid fraction (%)
4.12 MPa, 22 °C	70
2.8 MPa, – 40 °C	90

In order to examine the dispersion of CF<sub>3</sub>I in a cold environment, a new experimental facility will be designed and built. The experimental apparatus, the test matrix, and the test procedure to conduct the proposed discharge tests at sub-zero temperature are described in detail below.

## PROPOSED EXPERIMENTAL APPARATUS AND TEST PROCEDURE

The proposed experimental apparatus will be a simulated engine nacelle with airflow and with an annulus cross-sectional area and length scalable to a real nacelle. Figure 3 is a schematic of the proposed experimental fixture, which consists of a compartment with baffles, an agent release port, observation windows, and sampling ports. The compartment has a configuration and dimensions commensurate with a typical engine nacelle. The annulus has an inside diameter of  $\sim 0.5$  m and an outside diameter of  $\sim 1$  m, resulting in a cross-sectional area of  $\sim 0.59$  m<sup>2</sup>. The length ( $\sim 2$  m) of the compartment is comparable to the distance between the agent injection port and the downstream end of a typical engine nacelle. The compartment is made of stainless steel sheet metal.

The entire test facility will be placed inside a large environmental chamber. Agent discharge experiments will be conducted at the lowest achievable temperature (-40 °C) in the environmental chamber. Thermocouples will be placed on several locations inside the simulator to ensure thermal equilibrium with the compartment environment before a test is performed. Air flow through the compartment will be provided by a variable-speed blower.

The observations of the extinction of several tell-tale gaseous jet flames placed strategically at several locations in the compartment with video cameras will be used to assess *qualitatively* how well the agent is dispersed. These flames, which will be lit (possibly remotely) immediately before the discharge of  $CF_3I$ , are used as markers to detect the arrival of the agent to these locations. Extinction of a flame indicates the presence of sufficient agent concentration at that location. In addition, samples will be collected for  $\sim 0.5$  s immediately after agent discharge at locations close to the tell-tale flames using evacuated sample cylinders and subsequently analyzed for  $CF_3I$  using an FTIR. The integrated temporal  $CF_3I$  concentration measurements will be used to evaluate the uniformity of the agent dispersion in the compartment.

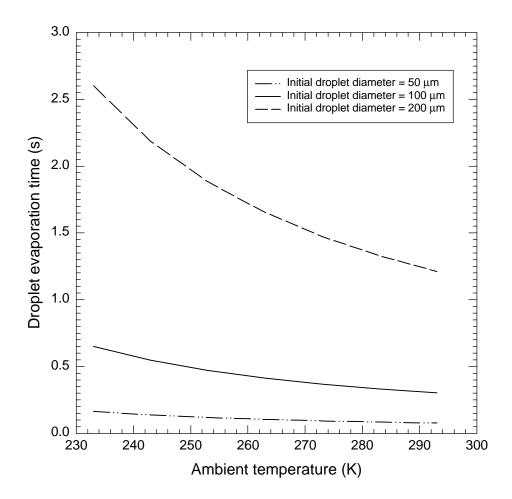


Figure 1. Evaporation time of a CF<sub>3</sub>I droplet as a function of ambient temperature when the ambient agent mass fraction is zero.

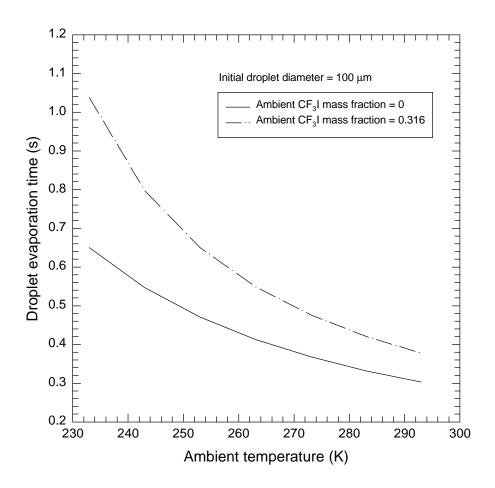


Figure 2. Effect of ambient agent mass fraction on the evaporation time of a 100  $\mu$ m CF<sub>3</sub>I droplet at various ambient temperatures.

Experiments using pure nitrogen will also be conducted to separate the effect of the pressurization gas (nitrogen) on flame extinction alone. If the flames cannot be extinguished by pure nitrogen, then one can infer that the extinguishment of these small flames is due to the presence of the agent. Baseline suppression performance will also be established using a room temperature environment.

Given the simulated nacelle volume, the amount of agent required for a fixed injection period ( $\sim 0.5$  s for typical nacelle applications) can be estimated using the generic nacelle modeling results in Gann (1995). The preparation of CF<sub>3</sub>I/dissolved nitrogen mixtures follows the procedure as described in Gann (1995). The release of the agent can be achieved using one of the several mechanisms (e.g., a Marotta valve or a squib). Agent injection time will be tailored using an orifice plate at the exit of the valve. Table 2 lists the proposed experimental matrix. Two representative air flows ( $\sim 1 \text{ kg/s}$  and  $\sim 0.3 \text{ kg/s}$ ) will be used in the tests.

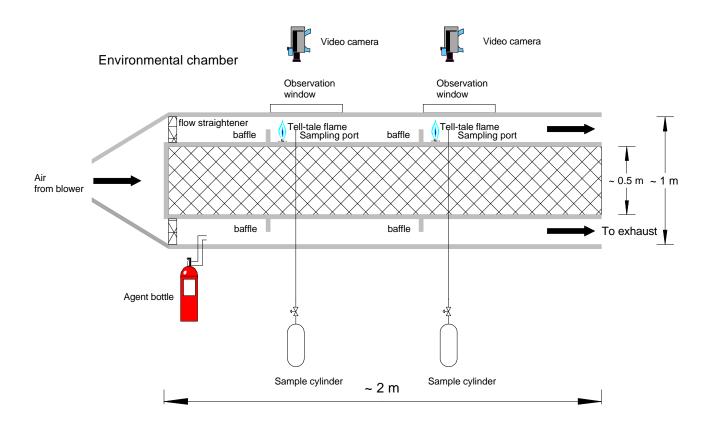


Figure 3. Proposed experimental apparatus for cold temperature agent dispersion.

Table 2 Proposed experimental matrix for cold temperature agent dispersion

Initial conditions of vessel	Conditions of vessel before	Conditions in simulated fire				
(half filled)	discharge	compartment				
22 °C and 4.12 MPa	− 40 °C at prevailing P <sup>¶</sup>	22 °C				
22 °C and 4.12 MPa	− 40 °C at prevailing P <sup>§</sup>	−40 °C §				
22 °C and 4.12 MPa	22 °C and 4.12 MPa	22 °C (baseline)				
22 °C and 2.75 MPa	− 40 °C at prevailing P <sup>¶</sup>	22 °C				
22 °C and 2.75 MPa	− 40 °C at prevailing P <sup>§</sup>	−40 °C §				
22 °C and 2.75 MPa	22 °C and 2.75 MPa	22 °C (baseline)				
Dry ign will be used to abill the vessel						

<sup>&</sup>lt;sup>1</sup>Dry ice will be used to chill the vessel.

The deliverables of the proposed research will be the performance data on  $CF_3I$  dispersion in a clustered compartment at -40 °C. The results from the  $CF_3I$  dispersion tests under -40 °C will determine if  $CF_3I$  can be effectively dispersed in an engine nacelle under temperature well below its normal boiling point.

<sup>§</sup> Tests will be performed in an environmental chamber.

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