in which ρ_s is the density of the solid particles.

To obtain the quantity $(1/u_s)(du_s/dx)$, we can measure the curvature of the particle streamline S(x) since

$$\frac{1}{u_s}\frac{du_s}{dx} = -\frac{S''(x)}{S'(x)}$$

Another way to obtain $(1/u_s)(du_s/dx)$ is by measuring the particle number density behind the shock n_s . If the number density is measured along a horizontal line x', then

$$\frac{1}{u_s}\frac{du_s}{dx} = -\frac{1}{n_s\sin\theta}\frac{\partial n_s}{\partial x'}$$

where θ is the shock angle. u_*/u is known exactly at the shock. By varying the wedge angle, the drag coefficients can be obtained over a range of relative Reynolds numbers.

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Velocity Profile in the Half-Jet Mixing Region of Turbulent Jets

Wing T. Chu*
University of Toronto, Toronto, Ontario, Canada

A WIDELY used similarity parameter for the nondimensional velocity U/U_0 of jets with exit velocity U_0 is the quantity $\sigma(Y-Y_{0.5})/x$. The x-axis is chosen parallel to the freestream direction or along the axis of an axisymmetric jet.

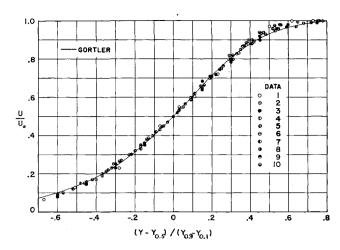


Fig. 1 Dimensionless velocity profile in the half-jet mixing region.

 $(Y-Y_{0.5})$ is the lateral distance between point of measurement and the point at which the velocity is half that of the exit velocity, and σ is a constant determined by fitting the measured velocity profile to the theoretical one of either Tollmein or Gortler. The quantity σ is not a universal constant and its value depends on the characteristics of an individual jet. The effect of Mach number on σ has been summed up in Fig. 12 of Ref. 1. See also Refs. 2 and 3. Because of the fairly wide scattering of data, it is hard to give a definite relationship between the two.

However, the variation of σ with different jet parameters (e.g., exit Reynolds number, Mach number, temperature, turbulence level) will be taken care of automatically if one uses the alternate parameter $(Y - Y_{0.5})/(Y_{0.9} - Y_{0.1})$ where $(Y_{0.9} - Y_{0.1})$ is the distance between the points at which the velocity is, respectively, 0.9 and 0.1 of the exit velocity $(U/U_0 = 0.9)$ and $U/U_0 = 0.1$. Abramovich⁴ used this parameter for the dimensionless velocity profile of his round jet and obtained a profile which fits well with that obtained from results of the plane jet of Albertson et al.⁵ Results from our own hot-wire measurements in a 4-in. low-speed round jet also agree well with Abramovich's profile. In view of this good agreement, a few experimental data of other investigators were compiled and replotted for comparison in Table 1 and Fig. 1.

It can be seen that, over a fairly wide range of exit velocities, the different velocity profiles lie very closely along the same curve. This universal curve is best fitted by Gortler's theoretical profile for incompressible jets⁶ with his parameter

Table 1 A list of some of the experimental investigations of turbulent jets

Investigation	Date	$egin{array}{c} \operatorname{Nozzle}^a \ & \operatorname{exit} \ & \operatorname{shape} \end{array}$	$egin{array}{c} \mathbf{Nozzle} \\ \mathbf{exit} \\ \mathbf{dimensions} \end{array}$	Axial distance from exit	Measurement method	$egin{array}{l} \mathbf{Nozzle} \ \mathbf{exit} \ \mathbf{speed} \end{array}$
Abramovich ⁴	1948	\overline{A}	100 mm	X/D = 2.5		40 m/sec
Liepmann and Laufer ²	1947	R	$60 \times 7\frac{1}{2}$ in.	$20~\mathrm{cm}$	Hot wire	59 fps
Laurence ³	1956	A	$3\frac{1}{2}$ in.	X/D = 4	Pitot pressure	M = 0.7
Davies & Fisher ⁷	1963	\boldsymbol{A}	1 in.	X/D = 3	Hot wire	M = 0.3
Bradshaw et al.8	1963	A	2 in.	X/D = 2	Hot wire	M = 0.3
Maydew and Reed ¹	1963	A	3 in.	X/D = 3	Pitot pressure	M = 0.95
Maydew and Reed ¹	1963	A	3 in.	X/D = 3	Pitot pressure	M = 1.49
Maydew and Reed ¹	1963	A	3 in.	X/D = 3	Pitot pressure	M = 1.96
Present investigation	1964	A	4 in.	X/D = 3	$\operatorname{Hot} \hat{\operatorname{Wire}}$	140 fps
Present investigation	1964	\boldsymbol{A}	4 in.	X/D = 4	Hot Wire	$140~\mathrm{fps}$

 $^{^{\}alpha}A$ = axisymmetric, R = rectangular.

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^{*} Graduate Student and Research Assistant, Institute for Aerospace Studies.

 $\sigma(Y - Y_{0.5})/X$ replaced by $(1.78)(Y - Y_{0.5})/(Y_{0.9} - Y_{0.1})$. The velocity profile in the mixing region of any particular jet can be estimated from the universal curve by determining the distances $Y_{0.1}$, $Y_{0.5}$, and $Y_{0.9}$ at one downstream location. [If these are not known, the line $Y_{0.5}$ vs X may be estimated and the relation

$$Y_{0.9} - Y_{0.1} = 1.78X/\sigma$$

resorted to, using tabulated estimates of σ where available (e.g., Refs. 1 and 2); this is in effect a reversion to the $\sigma(Y-Y_{0.5})/x$ similarity parameter and is only as accurate as the value of σ .

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Catalytic Probe Response to High Atom Flux in a Glow Discharge Shock Tube

W. P. Thompson* and R. A. Hartunian† Aerospace Corporation, Los Angeles, Calif.

IN Ref. 1 we reported on our measurements of heat transfer to a differential catalytic heat gage in shock-tube dissociated flows. The differential heat transfer is proportional to the atom concentration in the flow. Although most of the results followed generally expected trends, a new disturbing feature appeared in some of the measurements. Specifically, at the lower initial pressures in the shock tube, the differential heat transfer was found to vary with time instead of achieving a constant value immediately. At that time, we could envisage this effect as being due to either of two possibilities: first, slow dissociation relaxation in the shock layer of our probe, or secondly, a slow surface reaction rate at these lower pressures. We had pointed out at other times, and there has been some experimental evidence to the effect, that this latter possibility could be ascribed to submechanisms in the phenomenon of surface catalyzed atom recombination. example, an atom must be adsorbed onto a surface, it must diffuse to an active site where it awaits another atom to recombine with, and then the molecule desorbs. If the time scale for all of these events to occur is of the same order as the delivery rate of atoms in an experiment, it is conceivable that the observed time dependence of the differential catalytic heat-transfer measurements could be reconciled. That is, it is conceivable that the catalytic efficiency of a surface might depend on the atom flux. In our glow-discharge experiments,2 which are used to determine the catalytic efficiency of surfaces of interest, atoms are delivered to the surface at a relatively slow rate (1017/sec), whereas in our shock-tube experiments they are delivered on the order of 10²⁰/sec. Shock-tunnel flows will deliver atoms to the surface at a rate of about $5 \times 10^{19}/\text{sec}$. Since one of the major intended applications for our catalytic probe is to measure atom concentrations in hypersonic shock-tunnel test sections³ in order to establish the nonequilibrum state of the gas flow, it was imperative that we identify the source of the time dependence of our shock-tube measurements. It was clear that we needed an experiment in which a known step function of atoms be delivered to our catalytic surface at a rate of the order 10¹⁹/sec with a rise time of order ½ msec or less. Furthermore, it was essential that no gas phase kinetics be present so as to remove that possibility from causing a time dependent reading of our catalytic probe in these new experiments. To solve this problem we conceived the idea of the glow-discharge shock tube (GDST). The details of the construction and method of operation are given in a technical report⁴ just published at the Aerospace Corporation. Briefly, the GDST combines an ordinary glow-discharge tube and the shock tube. It is shown schematically in Fig. 1 together with an x-t diagram. The GDST works as follows: first, gas is flowed continuously along the glass tube (4 in. diam, 17 ft long, super-tough pyrex) at 10–30 fps, then, at a given time, the rf energy is suddenly applied at the position indicated by the coil in the sketch, which partially dissociates the gas; when the step function of dissociated gas so produced convects down to within a foot or so of the probe, the diaphragm of the driver section is made to burst with a solenoid-driven plunger, causing a shock to propagate into the predissociated gas. For the weak shocks that we employ $(M_s \bowtie 2)$, neither do the atoms recombine as a result of the increased density nor are any new atoms formed by thermal dissociation. Rather the only effect is to compress and accelerate to high speed the atoms already present in the tube. In our ordinary glow-discharge experiments, despite the fact that we suddenly turn the rf discharge on, we do not get a perfect step function of atoms, because as they convect down the tube at 20 fps, say, the atoms diffuse forward into the undissociated gas producing an atom profile that extends about 2 ft by the time it reaches the probe position. Under the slow flow, the front takes about 100 msec to cross the probe. On the other hand, after they are shocked, two particles on either end of this front cross our probe in about $\frac{1}{2}$ msec. This sharpening of the atom front, which can be seen on the x-t diagram, is one of the major effects sought. We have conducted three types of tests using an initial pressure of $600~\mu$ oxygen and a shock Mach number of 2. In the first, we

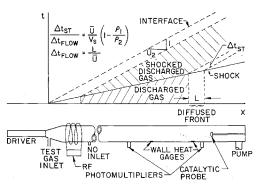


Fig. 1 Glow discharge shock tube; \overline{U} = convective velocity in slow flow; L = length of diffused front in slowflow; $V_s = \text{shock velocity and } \rho_2/\rho_1 = \text{shock density ratio.}$

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Member of Technical Staff, Aerophysics Department.

[†] Head, Aerophysics Department. Member AIAA.