

12

SECURITY CLASSIFICATION OF THIS PAGE (When Data Entered)

REPORT DOCUMENTATION PAGE		READ INSTRUCTIONS BEFORE COMPLETING FORM
1. REPORT NUMBER 14	2. GOVT ACCESSION NO. AD-A222549	3. RECIPIENT'S CATALOG NUMBER
4. TITLE (and Subtitle) Raman Investigation Of The Effects On Fused Silica Fibers		5. TYPE OF REPORT & PERIOD COVERED Technical Report #14
		6. PERFORMING ORG. REPORT NUMBER
7. AUTHOR(s) M.S. Hokmabadi and G.E. Walrafen		8. CONTRACT OR GRANT NUMBER(s) N00014-80-C-0305
9. PERFORMING ORGANIZATION NAME AND ADDRESS Department of Chemistry Howard University Washington, D.C. 20059		10. PROGRAM ELEMENT, PROJECT, TASK AREA & WORK UNIT NUMBERS NR-051-733
11. CONTROLLING OFFICE NAME AND ADDRESS Office of Naval Research Department of the Navy Arlington, Virginia 22217		12. REPORT DATE January 12, 1983
		13. NUMBER OF PAGES 8
14. MONITORING AGENCY NAME & ADDRESS (if different from Controlling Office)		15. SECURITY CLASS. (of this report) Unclassified
		15a. DECLASSIFICATION/DOWNGRADING SCHEDULE
16. DISTRIBUTION STATEMENT (of this Report) Approved for public release; reproduction is permitted for any purpose of the United States government distribution is unlimited.		
17. DISTRIBUTION STATEMENT (of the abstract entered in Block 20, if different from Report) Distributuon of this document is unlimited		
18. SUPPLEMENTARY NOTES Prepared and accepted for publication in the Journal of Chemical Physics.		
19. KEY WORDS (Continue on reverse side if necessary and identify by block number) Optical Fibers, Torsion, Raman Spectra		
20. ABSTRACT (Continue on reverse side if necessary and identify by block number) 7 Raman spectra have been obtained from fused silica optical fibers under high reversible torsion. These results are compared to Raman results from fused silica, irreversibly compacted to 90 kbar. An interpretation is presented which suggests a negative correlation between the mean Si-O-Si bridging angle and the mean Si-O distance, plus a negative correlation between the mean Si-O-Si bridging angle and the mean tilt angle between SiO ₄ tetrahedra.		

ADA 123549

DTIC FILE COPY

**"Raman Investigation Of The Effects Of Torsion On
Fused Silica Optical Fibers"**

by

**M.S. Hokmabadi and G.E. Walrafen
Chemistry Department
Howard University
Washington, D.C. 20059**



Codes		
and/or		
Special		
A		

The application of reversible tensile stress (3.3 GPa) to a pure fused silica optical fiber has been found to produce a significant modification of the Raman spectrum.⁽¹⁾ Other Raman changes due to irreversible uniaxial compaction (9 GPa),⁽²⁾ and reversible hydrostatic compression (1 GPa),⁽³⁾ were uncovered recently. The present work constitutes a continuation of such studies in which Raman changes due to reversible torsion on a fused silica optical fiber (at constant load) are described and compared to new Raman results⁽⁴⁾ from irreversibly compacted fused silica.

The application of reversible torsion to a horizontal fiber involved twisting one end, while the other end was rotationally immobilized just after passing over a vertical pulley. A constant downward force (load) was exerted by a weight attached to the fiber below the pulley. A 3.7 m section of a silicone clad pure fused silica fiber (4.5 m total length) was twisted 515 times before failure,⁽⁵⁾ corresponding to 501 degree/cm or 8.75 radian/cm. The resulting spectrum is shown in fig. 1(b).

Comparison of the untwisted fiber spectrum, 1(a), to that of the twisted fiber, 1(b), indicates that one significant Raman change due to torsion involves a loss of intensity between 0-450 cm^{-1} . This result is evident from the $\sim 16 \text{ cm}^{-1}$ half-width decrease shown, and by the difference spectrum (inset, upper left).

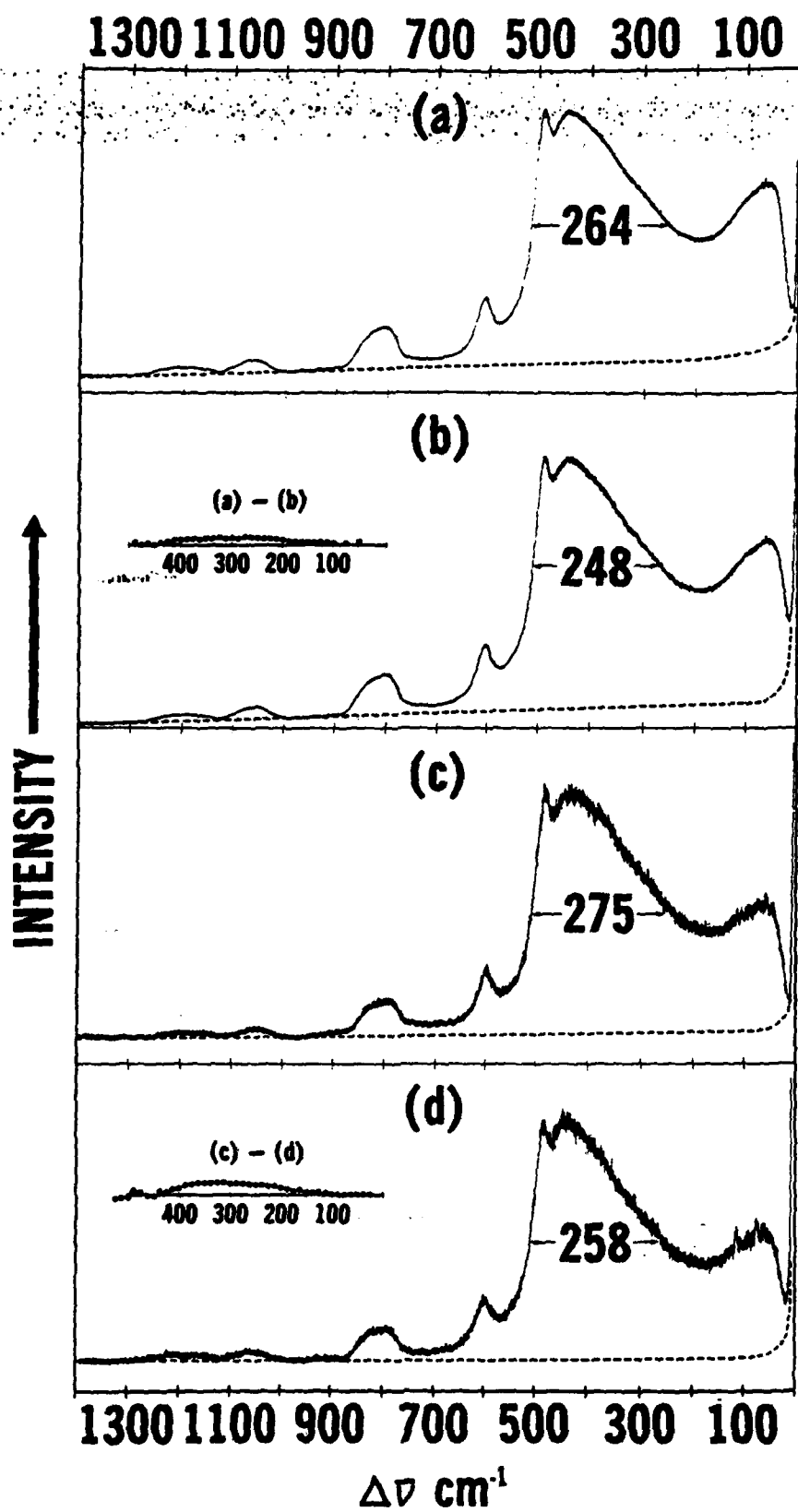
Other Raman modifications similar to those of torsion are evident from spectra of uncompacted (parent) and irreversibly compacted (9 GPa) fused silica, figs. 1(c) and 1(d), respectively.^(2,4) Here, the half-width decreases $\sim 17 \text{ cm}^{-1}$, and the difference spectrum (inset, lower left) is also similar to the one above.

Close examination of the two difference spectra, however, indicates that they are not quite identical. The lower spectrum peaks at a higher frequency, $\sim 350 \text{ cm}^{-1}$, and a decrease in the intensity of the "defect" peak⁽⁶⁾ at 490 cm^{-1} is apparent—an effect not previously detected under lower S/N.⁽²⁾ But measurements involving several spectra from twisted and untwisted fibers show that a similar, but smaller, intensity decrease at 490 cm^{-1} also occurs.

From all of the Raman spectral modifications examined for reversible tensile stress,⁽¹⁾ irreversible uniaxial compaction,⁽²⁾ reversible hydrostatic compression,⁽³⁾ and irreversible neutron compaction⁽⁷⁾ it appears that densification decreases the Raman intensity between 0-450 cm^{-1} . Generalizations including the 490 and 605 cm^{-1} "defect" peaks are harder to make. But an increase of intensity between 0-450 cm^{-1} , and an intensity increase at 490 cm^{-1} were observed for reversible tensile stress, whereas just the opposite effects are now

CAPTION

Figure 1. Raman spectra from untwisted (a) and twisted (b) pure fused silica optical fibers, and from (parent) uncompacted (c) and compacted (9 GPa) bulk fused silica (d). The change in half-width from (a) to (b), 16 cm^{-1} , agrees favorably with the change in half-width from (c) to (d), 17 cm^{-1} , but the (a) and (b) half-widths should not be compared to the (c) and (d) half-widths because different samples and conditions were involved.



found for reversible torsion and irreversible compaction.

Raman intensity changes between 0-450 cm^{-1} , if large, also entail frequency shifts at 800 cm^{-1} and 1060 cm^{-1} . These effects may result from correlation between the mean tilt angle between SiO_4 tetrahedra, $\bar{\delta}$, the mean Si-O-Si bridging angle, $\bar{\phi}$, and the mean Si-O bond length, $\bar{r}$. For amorphous silica only, we suggest that an increase of $\bar{\delta}$ refers to a decrease of $\bar{\phi}$ and an increase of $\bar{r}$ (negative correlation between $\bar{\phi}$ and $\bar{r}$ is well-established for fused silica, (8) and a negative correlation between $\bar{\delta}$ and $\bar{\phi}$ has been reported for pressurized α -quartz. (9)) The Raman intensity between 0-450 cm^{-1} is further suggested to decrease as $\bar{\delta}$ increases.

For reversible hydrostatic compression (3) and for irreversible neutron compaction, (7) a large loss of Raman intensity between 0-450 cm^{-1} is accompanied by an upward frequency shift at 800 cm^{-1} and by a downward frequency shift at 1060 cm^{-1} . With decreasing ϕ , an increase in the Si-O-Si bending force constant is thought to move the 800 cm^{-1} peak upward in frequency, (7) and a decrease in the Si-O stretching force constant due to an increase in r is thought to move the 1060 cm^{-1} peak position downward.

From the correlations now suggested, namely, that increasing $\bar{\delta}$ refers to a decrease in the Raman intensity between 0-450 cm^{-1} , I_{0-450} , as well as an upward shift at 800 cm^{-1} (increasing $\Delta\bar{\nu}$) and a downward shift at 1060 cm^{-1} (decreasing $\Delta\bar{\nu}$), a table may be constructed involving tension, the three compressions, and torsion. Missing entries refer to frequency shifts at 800 or 1060 cm^{-1} which are presently undetectable because they scale with the small Raman intensity changes between 0-450 cm^{-1} . Nevertheless, this table suggests that $\bar{\delta}$ decreases with tensile stress, (1) whereas $\bar{\delta}$ is suggested to increase for the three compressive cases, (2), (3), (7) and also for reversible torsion.

A more detailed interpretation involving Gaussian deconvolution (11) of the 0-450 cm^{-1} region will appear subsequently.

Dr. P. N. Krishnan is thanked for helpful suggestions. This work was supported by contracts from the Office of Naval Research.

REFERENCES

- (1) G. E. Walrafen, P. N. Krishnan, and S. W. Freiman, J. Appl. Phys. 52, 2832 (1981).
- (2) G. E. Walrafen and P. N. Krishnan, J. Chem. Phys. 74, 5328 (1981).
- (3) M. S. Hokmabadi, Doctoral Dissertation, Howard University, 1981.
- (4) Scrambled laser polarization was used prior to Raman excitation, Figs. 1(c) and 1(d), to counteract effects of inhomogeneities (cracks) in the compacted sample.
- (5) Further rotation caused a uniform pulverization of the fiber over the entire twisted region.
- (6) The "defect" peaks at 490 and 605 cm⁻¹ have recently been re-assigned, A. K. Revesz and G. E. Walrafen, J. Noncryst. Solids (in press).
- (7) R. H. Stolen, J. T. Krause and C. R. Kurkjian, Discuss. Faraday Soc. 50, 103 (1970); and, J. B. Bates, R. W. Hendricks, and L. B. Shaffer, J. Chem. Phys. 61, 4163 (1974).
- (8) G. V. Gibbs, E. P. Meagher, M. D. Newton and P. K. Swanson in Structure and Bonding in Crystals, Vol. 1 (M. O'Keefe et al., eds) p. 195, Academic Press, New York (1981).
- (9) J. D. Jorgensen, J. Appl. Phys. 49, 5473 (1978).
- (10) G. E. Walrafen, M. Abebe, F. A. Mauer, S. Block, G. J. Piermarini and R. G. Munro, J. Chem. Phys. 77, 2166 (1982).
- (11) G. E. Walrafen and P. N. Krishnan, Appl. Optics 21, 359 (1982).

TECHNICAL REPORT DISTRIBUTION LIST, GEN

	<u>No. Copies</u>		<u>No. Copies</u>
Office of Naval Research Attn: Code 413 800 North Quincy Street Arlington, Virginia 22217	2	Naval Ocean Systems Center Attn: Mr. Joe McCartney San Diego, California 92152	1
ONR Pasadena Detachment Attn: Dr. R. J. Marcus 1030 East Green Street Pasadena, California 91106	1	Naval Weapons Center Attn: Dr. A. B. Amster, Chemistry Division China Lake, California 93555	1
Commander, Naval Air Systems Command Attn: Code 310C (H. Rosenwasser) Department of the Navy Washington, D.C. 20360	1	Naval Civil Engineering Laboratory Attn: Dr. R. W. Drisko Port Hueneme, California 93401	1
Defense Technical Information Center Building 5, Cameron Station Alexandria, Virginia 22314	12	Dean William Tolles Naval Postgraduate School Monterey, California 93940	1
Dr. Fred Saalfeld Chemistry Division, Code 6100 Naval Research Laboratory Washington, D.C. 20375	1	Scientific Advisor Commandant of the Marine Corps (Code RD-1) Washington, D.C. 20380	1
U.S. Army Research Office Attn: CRD-AA-IP P. O. Box 12211 Research Triangle Park, N.C. 27709	1	Naval Ship Research and Development Center Attn: Dr. G. Bosmajian, Applied Chemistry Division Annapolis, Maryland 21401	1
Mr. Vincent Schaper DTNSRDC Code 2803 Annapolis, Maryland 21402	1	Mr. John Boyle Materials Branch Naval Ship Engineering Center Philadelphia, Pennsylvania 19112	1
Naval Ocean Systems Center Attn: Dr. S. Yamamoto Marine Sciences Division San Diego, California 91232	1	Mr. A. M. Anzalone Administrative Librarian PLASTEC/ARRADCOM Bldg 3401 Dover, New Jersey 07801	1

TECHNICAL REPORT DISTRIBUTION LIST, 051A

	<u>No. Copies</u>		<u>No. Copies</u>
Dr. M. A. El-Sayed Department of Chemistry University of California, Los Angeles Los Angeles, California 90024	1	Dr. M. Rauhut Chemical Research Division American Cyanamid Company Bound Brook, New Jersey 08805	1
Dr. E. R. Bernstein Department of Chemistry Colorado State University Fort Collins, Colorado 80521	1	Dr. J. I. Zink Department of Chemistry University of California, Los Angeles Los Angeles, California 90024	1
Dr. C. A. Heller Naval Weapons Code 6059 China Lake, California 93555	1	Dr. D. M. Burland IBM San Jose Research Center 5600 Cottle Road San Jose, California 95143	1
Dr. J. R. MacDonald Chemistry Division Naval Research Laboratory Code 6110 Washington, D.C. 20375	1	Dr. John Cooper Code 6130 Naval Research Laboratory Washington, D.C. 20375	1
Dr. G. B. Schuster Chemistry Department University of Illinois Urbana, Illinois 61801	1	Dr. William M Jackson Department of Chemistry Howard University Washington, D.C. 20059	1
Dr. A. Adamson Department of Chemistry University of Southern California Los Angeles, California 90007	1	Dr. George E. Walrafen Department of Chemistry Howard University Washington, D.C. 20059	1
Dr. M. S. Wrighton Department of Chemistry Massachusetts Institute of Technology Cambridge, Massachusetts 02139	1	Dr. Joe Brandelik AFWAL/AADO-1 Wright Patterson AFB Fairborn, Ohio 45433	1
Dr. A. Paul Schaap Department of Chemistry Wayne State University Detroit, Michigan 49207	1	Dr. Gary Bjorklund IBM 5600 Cottle Road San Jose, California 95143	1
		Dr. Carmen Ortiz Cousejo Superior de Investigaciones Cientificas Serrano 117 Madrid 6, Spain	1