



Designation: F1438 – 93 (Reapproved 2020)

Standard Test Method for Determination of Surface Roughness by Scanning Tunneling Microscopy for Gas Distribution System Components¹

This standard is issued under the fixed designation F1438; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ϵ) indicates an editorial change since the last revision or reapproval.

INTRODUCTION

Semiconductor clean rooms are serviced by high-purity gas distribution systems. This test method presents a procedure that may be applied for the evaluation of one or more components considered for use in such systems.

1. Scope

1.1 The purpose of this test method is to define a method for analyzing the surface texture of the above-mentioned components using a scanning tunneling microscope (STM). STM is a noncontact method of surface profiling that can measure three-dimensional surface features in the nanometer size range, which can then be used to represent the surface texture or to provide figures of merit. Application of this test method, where surface texture is used as a selection criterion, is expected to yield comparable data among different components tested.

1.2 Limitations:

1.2.1 This test method is limited to characterization of stainless steel surfaces that are smoother than $R_a = 0.25 \mu\text{m}$, as determined by a contact-stylus profilometer and defined by ANSI B46.1. The magnifications and height scales used in this test method were chosen with this smoothness in mind.

1.2.2 Intentional etching or conductive coating of the surface are considered modifications of the gas-wetted surface and are not covered by this test method.

1.2.3 This test method does not cover steels that have an oxide layer too thick to permit tunneling under the test conditions outlined in 11.3.

1.3 This technique is written with the assumption that the STM operator understands the use of the instrument, its governing principles, and any artifacts that can arise. Discussion of these points is beyond the scope of this test method.

1.4 The values stated in SI units are to be regarded as the standard.

1.5 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

1.6 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

2.1 ASTM Standards:

[E691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method](#)²

2.2 ANSI Standard:

[ANSI B.46.1-85, "Surface Texture \(Surface Roughness, Waviness, and Lay\)," ANSI/ASME, 1985](#)³

3. Terminology

3.1 Definitions of Terms Specific to This Standard:

3.1.1 *artifact*—any contribution to an image from other than true surface morphology. This could include such examples as vibration, electronic noise, thermal drift, or tip imperfections.

3.1.2 *center line (graphical center line)*—line parallel to the direction of profile measurement, such that the sum of the areas contained between it and the profile contained on either side are equal (see Calculation Section).

¹ This test method is under the jurisdiction of ASTM Committee F01 on Electronics and is the direct responsibility of Subcommittee F01.10 on Contamination Control.

Current edition approved April 15, 2020. Published May 2020. Originally approved in 1993. Last previous edition approved in 2012 as F1438–93(2012). DOI: 10.1520/F1438-93R20.

² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

³ Available from American National Standards Institute, 13th Floor, 11 W. 42nd St., New York, NY 10036.

3.1.3 *cutoff length* (l_c)—for profiles in this context, the sampling length, that is, the length of a single scan, in nanometers (see Calculation Section).

3.1.4 *current*—in this context, the tunneling current (expressed in nanoamperes) that flows in either direction between the tip and surface, under the conditions specified.

3.1.5 *feature height*—datum (height in the z-direction) of any point in the scan area, relative to the lowest point in the scan area, as derived from tunneling current during tip rastering.

3.1.6 *filter*—process of modification of surface data for purposes of numerical analysis or data presentation. Examples include high or low pass filters and plane-fitting.

3.1.7 *gold ruled grating*—gold surface having uniformly spaced grooves of known depth and separation; used for micrometer scale x-y calibration.

3.1.8 *illuminated surface*—three-dimensional image representation that simulates a reflective surface illuminated obliquely or from overhead.

3.1.9 *image*—surface topography represented by plotting feature height as a function of tip position. The feature height data is derived from the amount of tunneling current flowing between the tip and surface during rastering.

3.1.10 *line plot*—three-dimensional image given as side-by-side surface profiles.

3.1.11 *mean roughness* (R_a)—average deviation from the mean of all profile heights (see algebraic definition in the Calculation Section).

3.1.12 *peak*—highest point between two crossing points of a profile and its center line.

3.1.13 *profile*—the cross-sectional data that has been high pass filtered with a two-pole filter having a gain of 75 % at the cutoff length l_c (in nanometers).

3.1.14 *raster*—repetitive scanning in the x-direction while moving stepwise in the y-direction; also the area defined by such action.

3.1.15 *scan*—a single, continuous movement in one direction (defined as the x-direction) of the tip relative to sample surface.

3.1.16 *scan area*—area covered by successive, side by side scans.

3.1.17 *scan length*—distance from start to end of a single scan, without moving in the y-direction (see cutoff length).

3.1.18 *scan rate*—the speed at which the tip moves relative to the surface.

3.1.19 *shaded height plot*—image representing feature height as dark or light shades (any color) over a two-dimensional area. Higher features are shaded lighter and lower features are shaded darker.

3.1.20 *thermal drift*—movement of the surface with respect to the tip due to a lack of thermal equilibrium.

3.1.21 *tilted surface*—three-dimensional image showing surface tilted away from viewer, as opposed to a topview.

3.1.22 *tip crash*—touching of tip to surface, during rastering or attempts to initiate tunneling, usually resulting in damage to one or both.

3.1.23 *top view*—image represented as a surface viewed from overhead.

3.1.24 *tunneling*—in this context, the flow of current between the tip and surface (see **current**); more discussion can be found in additional references.⁴

3.1.25 *valley*—lowest point between two crossing points of a profile and its center line.

3.1.26 *voltage*—bias voltage, expressed in volts (V) or millivolts (mV), applied between the tip and the surface.

3.2 Abbreviations and Symbols:

3.2.1 *HOPG*—highly ordered pyrolytic graphite; used for atomic scale x-y calibration of the scanning tunneling microscopy.

3.2.2 *STM*—scanning tunneling microscopy (or microscope).

3.2.3 *nA*—nanoamperes (1×10^{-9} amperes).

3.2.4 *Pt/Ir*—platinum and iridium alloy wire used to make tunneling tips.

3.2.5 R_a —see mean roughness.

3.2.6 R_{max} —maximum height difference between the highest and the lowest points on the profile over the length of the profile (see Calculation Section).

3.2.7 *root mean square (RMS)*—see algebraic definition in Calculation Section.

3.2.8 R_z —the 10-point mean roughness; that is, the average difference in height between the five highest peaks and the five lowest valleys over the length of the profile (see Calculation Section).

3.2.9 *x-direction*—see *scan*.

3.2.10 *y-direction*—the direction, in the sample plane, over which successive scans are taken, orthogonal to the scan direction.

3.2.11 *z-direction*—the direction perpendicular to the sample plane. Also referred to as the feature height direction.

3.2.12 Z_t —same as feature height (see Calculation Section).

3.2.13 Z_{max} —maximum height difference over entire surface (see Calculation Section).

3.2.14 Z_{rms} —root-mean-square of all surface heights (see Calculation Section).

4. Summary of Test Method

4.1 In this test method a sharp, conductive tip is scanned over very closely but not in contact with a conductive surface; that is, they are separated by a gap of several angstroms. A bias voltage present between them causes a flow of electrons through, rather than over, the energy barrier represented by this tip-surface gap. This flow is referred to as the tunneling

⁴Binning, G., et al., "Surface Studies by Scanning Tunneling Microscopy," *Physical Review Letters*, Vol 49, No. 1, July 1982, pp. 57-61.

current. The manner in which the current fluctuates during the scanning process is used to indicate the surface's topography. Though the tip or sample can be scanned, the method described here considers only the tip to be in motion. A more extensive discussion of the operating principles can be found in other literature.

4.2 In this test method, stainless steel tubing is used as an example of a component surface. An area of the surface is first scanned at a width of 500 nA, then 2000 nA. Even though larger areas can be scanned by most instruments, these magnifications are chosen to show surface texture in a size range beyond that measured by contact stylus type surface profiling instruments, but not at an atomic scale. The surface scans are then compared for damage, artifacts, etc. Numerical analysis can then proceed using these data for roughness or surface area or both, following the model of other standards such as ANSI B46.1.

5. Significance and Use

5.1 The use of STM images and data is for purposes of textural quality assessment and calculation of figures of merit, and for high purity gas system clean room components.

5.2 This test method defines a standard data presentation format and suggests figures of merit that utilize STM's ability to analyze three-dimensional surface features.

6. Interferences

6.1 Some (stainless steel) component surfaces have an oxide layer that prevents tunneling from occurring under any conditions without affecting tip or surface morphology. This results in ambiguous surface data. Such surfaces require the use of other techniques for topographic measurement.

6.2 This test method assumes that the images obtained are unperturbed by very thin, non-solid layers (for example, hydrocarbons, moisture) on the surface.

6.3 Operation with the surface in air, vacuum, or under inert liquids is permissible. (The liquids must be suitably inert and fluid, so as to not modify the apparent surface topography or introduce artifacts into the image.) Water is not recommended.

6.4 The tip shall be made from platinum/iridium or tungsten.

NOTE 1—Caution: The tip must not have previously touched any surface (see 11.7).

7. Apparatus

7.1 *Scanning Tunneling Microscope*, capable of the following may be used:

- 7.1.1 Scanning lengths up to at least 50 μm ,
- 7.1.2 Sustaining ± 3 V between the tip and sample,
- 7.1.3 Monitoring tunneling current as low as 0.05 nA,
- 7.1.4 Traversing feature height variations as great as 2 μm without touching the tip to the surface, and
- 7.1.5 Providing the surface topography as a shaded height plot or line plot.

7.2 Inert atmospheres, temperature controls, acoustic isolation, and vibration isolation are to be provided as necessary to obtain artifact-free images.

8. Sampling

8.1 Many components are too large or irregularly shaped to permit STM analysis without cutting a sample from the component. Low speed cutting, preferably without lubricants, using a diamond blade saw is recommended over high speed abrasive cutting or hacksaws.

8.2 This sampling must not modify the surface topography, such as effects due to stress, heat, corrosion, or combination thereof, from its condition as found in the component.

8.3 Cleaning the surface using an inert fluid to remove cutting contamination is permitted.

9. Calibration

9.1 Calibration frequency may vary with different instrument manufacturers. It should be performed, at least initially, then yearly, and after any repair or addition to the instrument's hardware and software.

9.2 Following the manufacturer's recommendations, the STM will be calibrated using the HOPG, gold ruled grating, or some other suitable dimensional standard, depending upon the size range to be used and the accuracy of the standard.

10. Conditioning

10.1 A conductive path for the sample shall be provided for voltage biasing of the sample with respect to the tip.

10.2 Mount the sample so the tip will scan an arbitrarily chosen representative area.

10.3 Bring the sample and microscope to thermal equilibrium.

11. Procedure

11.1 As stated in 1.2.3, this test method does not cover steels that have an oxide layer too thick to permit tunneling under the test conditions outlined in this test method.

11.2 Make sure that a minimum of 200 data points is collected in the x-direction, and at least 200 scans per raster is in the y-direction ($200 \times 200 = 40\,000$ data points).

11.3 Initiate tunneling between a Pt/Ir tip and the sample in accordance with the manufacturer's instructions under conditions to provide artifact free images. Suggested starting values: voltage bias of 1800 mV, and current levels monitored at 0.1 nA (*Standard Test Conditions*—Room temperature and ambient pressure (101.3 kPa, $25^\circ \pm 2^\circ\text{C}$)).

11.4 Scan an area 500-nm across at a rate of approximately 2 s/ μm (or slow enough to prevent the tip from touching the surface during rastering), collecting at least 200 data points with each x-direction scan.

11.5 Collect area scans so that comparisons can be made between the first two successive rasters. Ensure that the area has not drifted more than 10 % of the scan width in any direction. If it has, discontinue scanning for sufficient time to

allow thermal equilibrium to be obtained. Determine this by repeating 11.3 through 11.5.

11.6 Scan the selected 500-nm area for at least five full, successive rasters and then store at least the last (fifth) image of that scan area. Fig. 1, Fig. 2, and Fig. 3 show an example of such an area after the fifth successive raster (see data presentation section for explanation of format shown).

11.7 Inconsistent topography from one scan to the next is a sign of tip crashing. Fig. 4 shows an example of where the tip touched the surface during rastering.

11.8 Change the scanned width to 2000 nm across, centered around the same region, and immediately obtain an image of that area. Fig. 5, Fig. 6, and Fig. 7 show an example of a 2000-nm wide area.

11.9 Observe whether the 2000-nm image shows evidence of damage from scanning the previous 500-nm (smaller) area. This generally appears as a 500-nm square of very different topography or height near the center of the 2000-nm image. If there is such evidence, the surface may be too oxidized to analyze (see 6.1). Fig. 8 shows an example of damage apparent at 2000 nm, which occurred during the 500 nm-wide rasters.

11.10 If no scanning damage is apparent, store the data for the 2000-nm area.

11.11 Disengage tunneling, relocate the tip or sample to a different surface area, and repeat 11.4 through 11.10 using the best imaging conditions. Make sure that the distance moved is at least far enough to prevent reanalyzing a portion of the previously scanned areas.

11.12 If there is any uncertainty about tip artifacts or tip-crashing, replace the tip and repeat 11.3 through 11.10.

11.13 Calculations of figures of merit (see Calculation Section) should utilize data from at least three different areas of the same scan width. If such figures are required, analyze additional areas in accordance with 11.3 through 11.10.

12. Calculations

12.1 Figures of merit can be calculated from the surface area or surface profiles data.

12.2 Surface data consists of successive scans across an area. Profiles extracted from the surface data that cross these scans may contain artifacts due to slight discontinuities from scan to scan, which may appear as small peaks in the profile. If these artifacts are present, then use only profiles taken in the scan direction.

12.3 The results of numerical analysis of the data may include the following, as defined in 12.10.1 and 12.10.2, in tabular form:

R_a , R_{max} , and R_z (from profiles).

Z_{max} , and Z_{rms} (from surface areas).

12.4 As stated in the profile definition, the surface profile (cross-sectional data) must be cutoff filtered before R_a or R_z can be determined.

12.5 Calculate profile figures of merit (R_a and R_z) from data for five profiles each on at least three different areas on the same sample surface (for a total of 15 profiles of a single scan

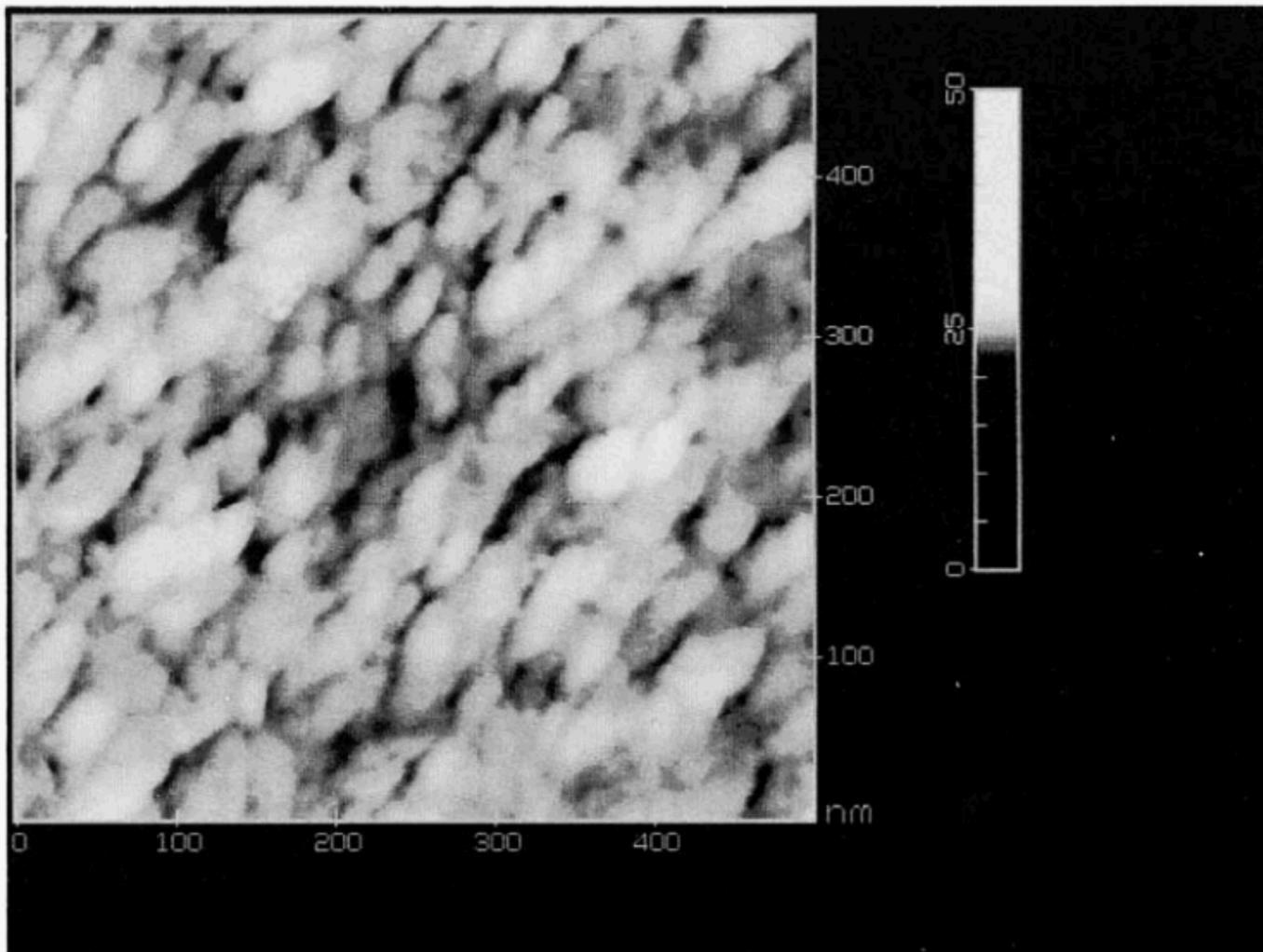


FIG. 1 Shaded Height Topview of Electropolished 316L Stainless Steel, 500-nm Across

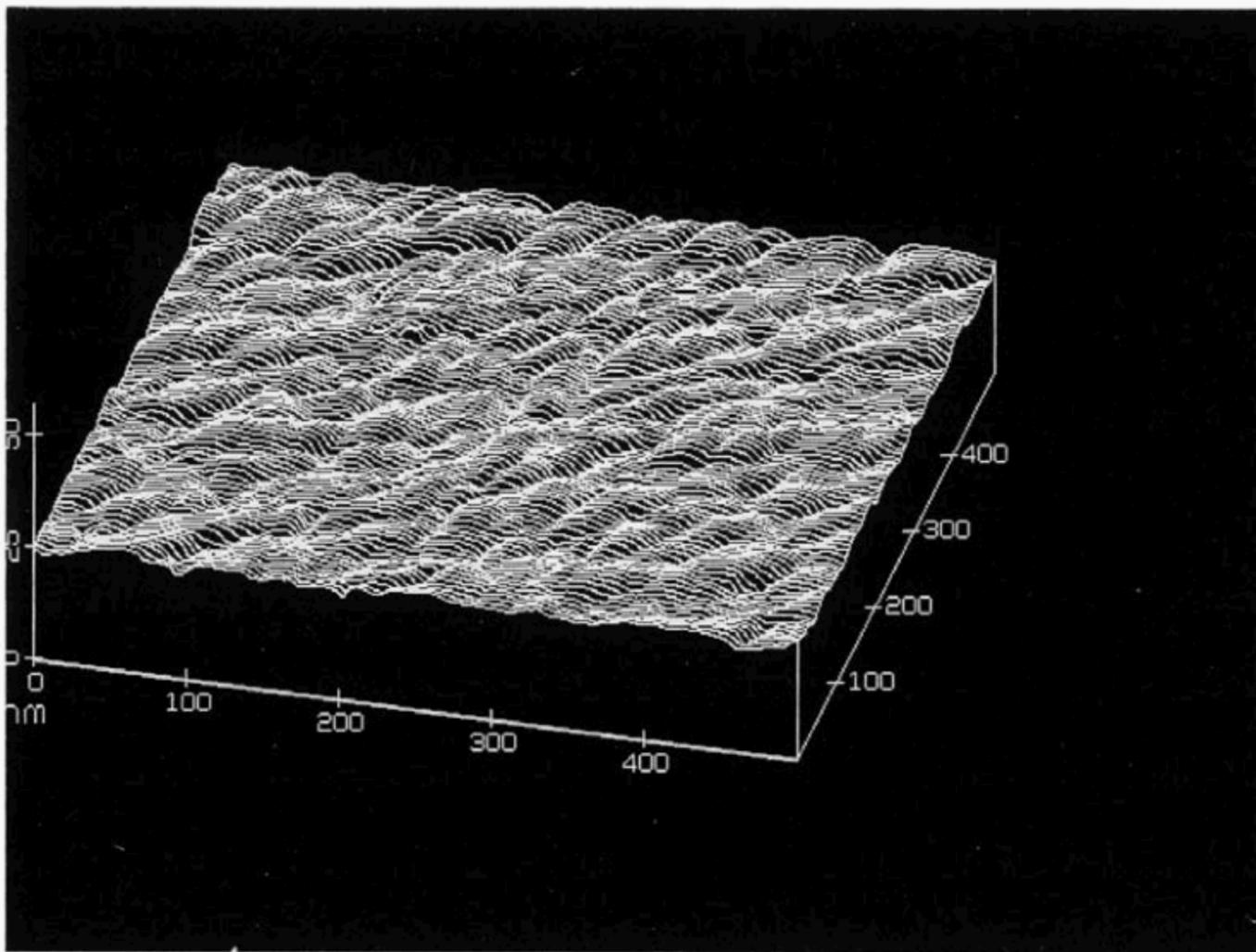


FIG. 2 Line Plot of Same Scan as Shown in Fig. 1

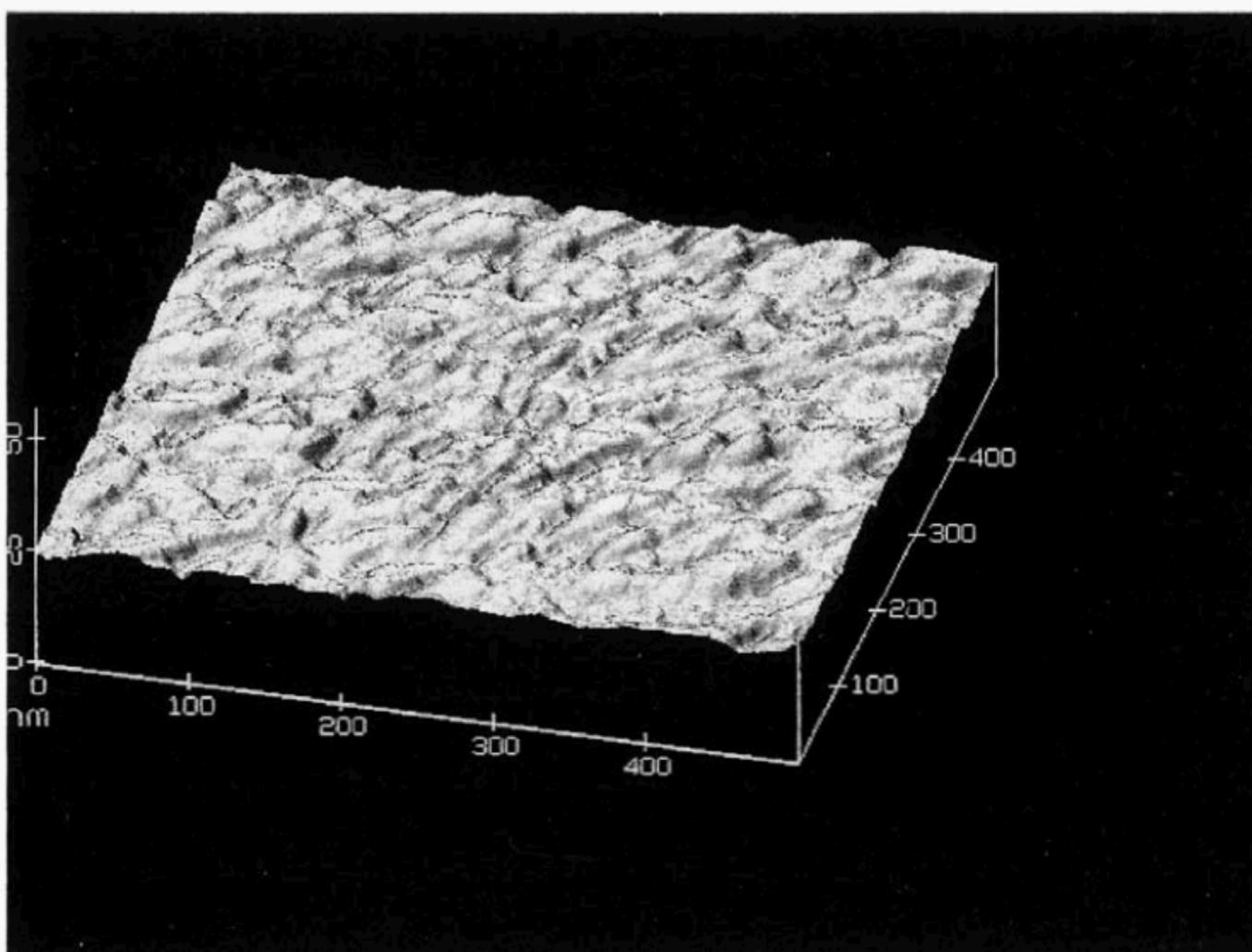


FIG. 3 Top-Illuminated (Tilted Surface) View of Same Scan as Shown in Fig. 1

length). Also report a 95 % confidence limit of the mean in the table, along with the profile length in nanometers.

12.6 Calculate area figures of merit (Z_{\max} and Z_{rms}) from data for at least three different areas on the same sample

surface. The average of at least three such areas will be the reported values in the table.

12.7 Table 1 shows results for the surfaces shown in Fig. 1, Fig. 5, and Fig. 9.

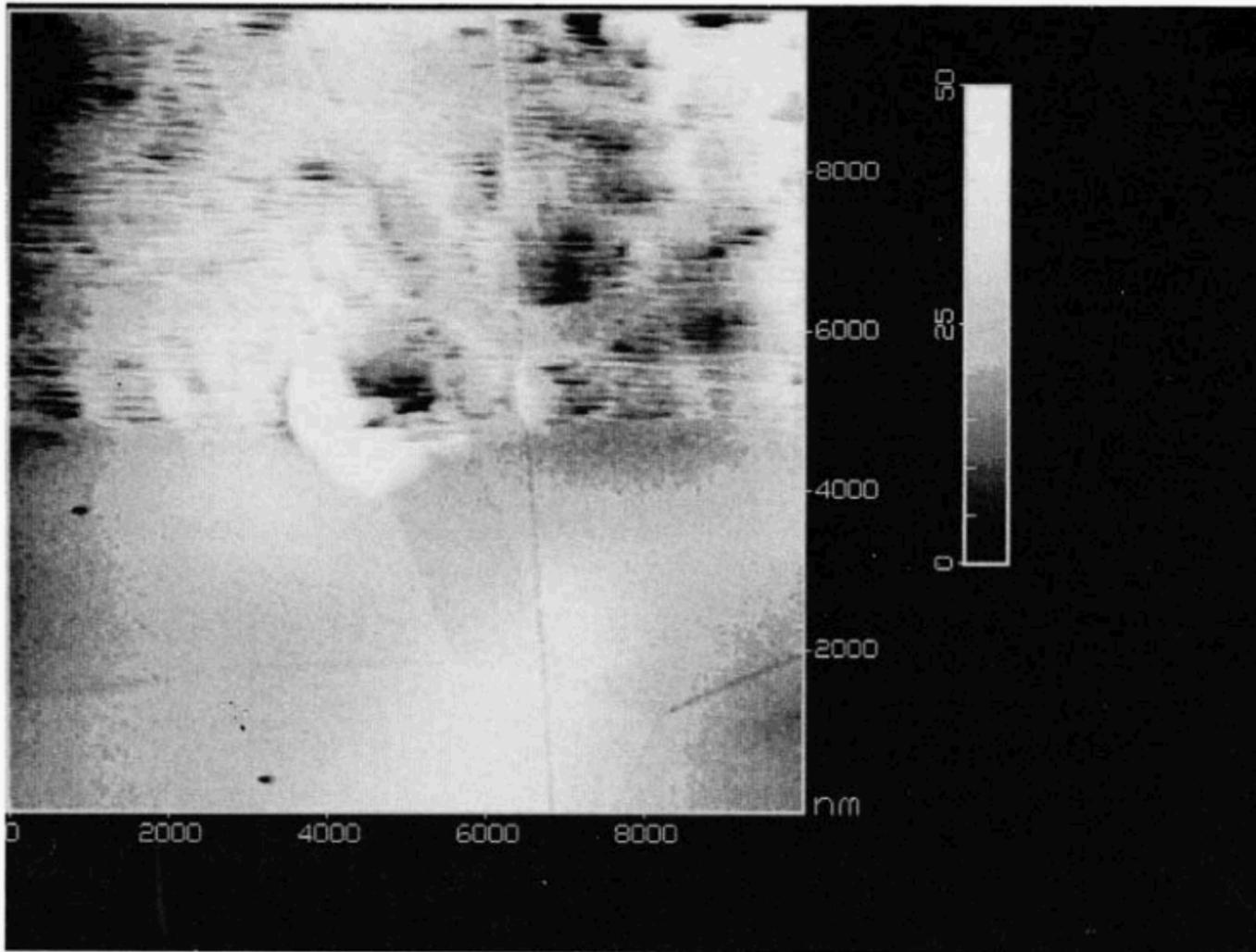


FIG. 4 Shaded Height of Topview of Stainless Steel Surface Showing Example of Tip “Crashing” Artifact (Upper Portion is Distorted Due to Damaged Tip.)

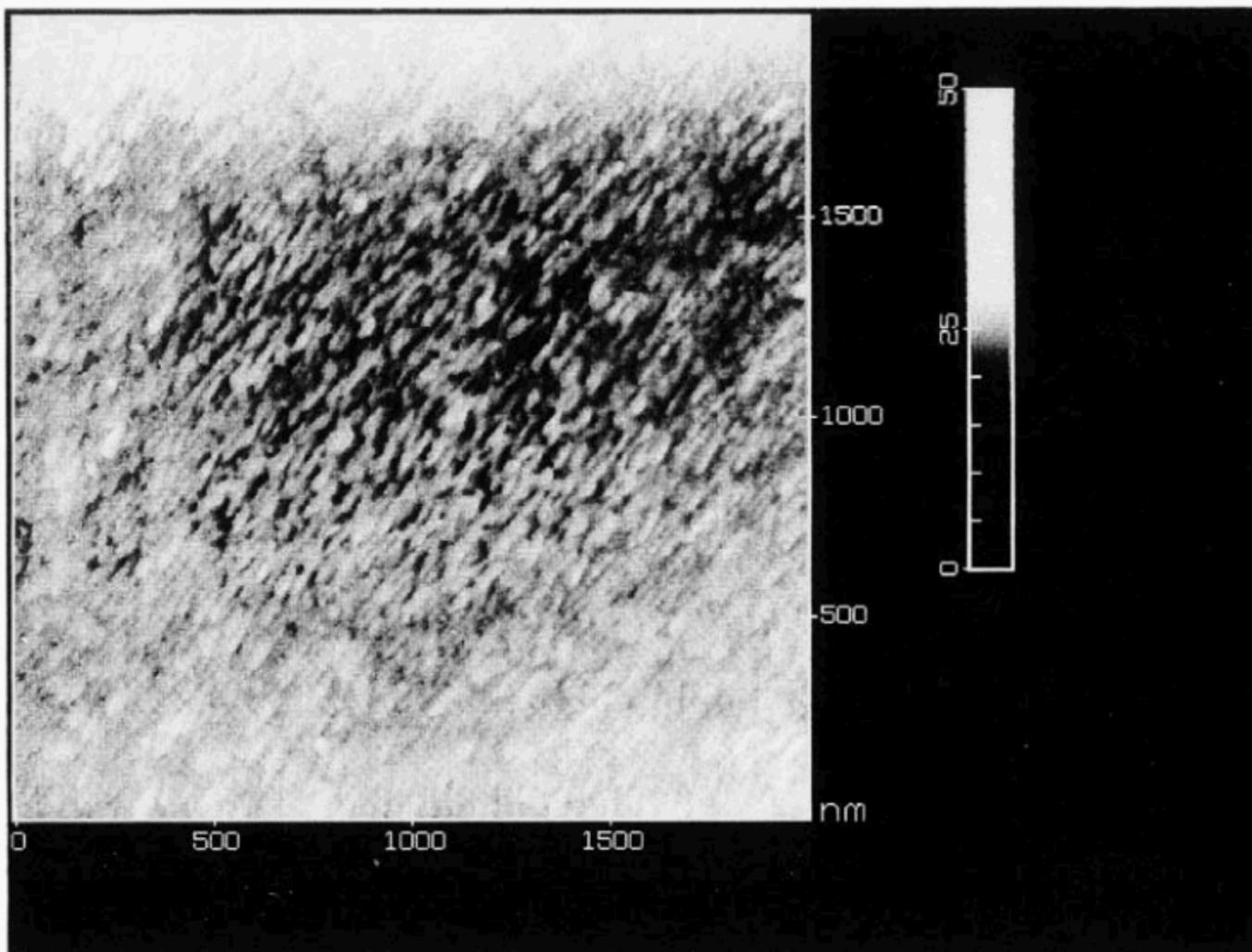


FIG. 5 Shaded Height Topview of Electropolished Stainless Steel, 2000-nm Across

12.8 The units for each will be reported in nanometers.

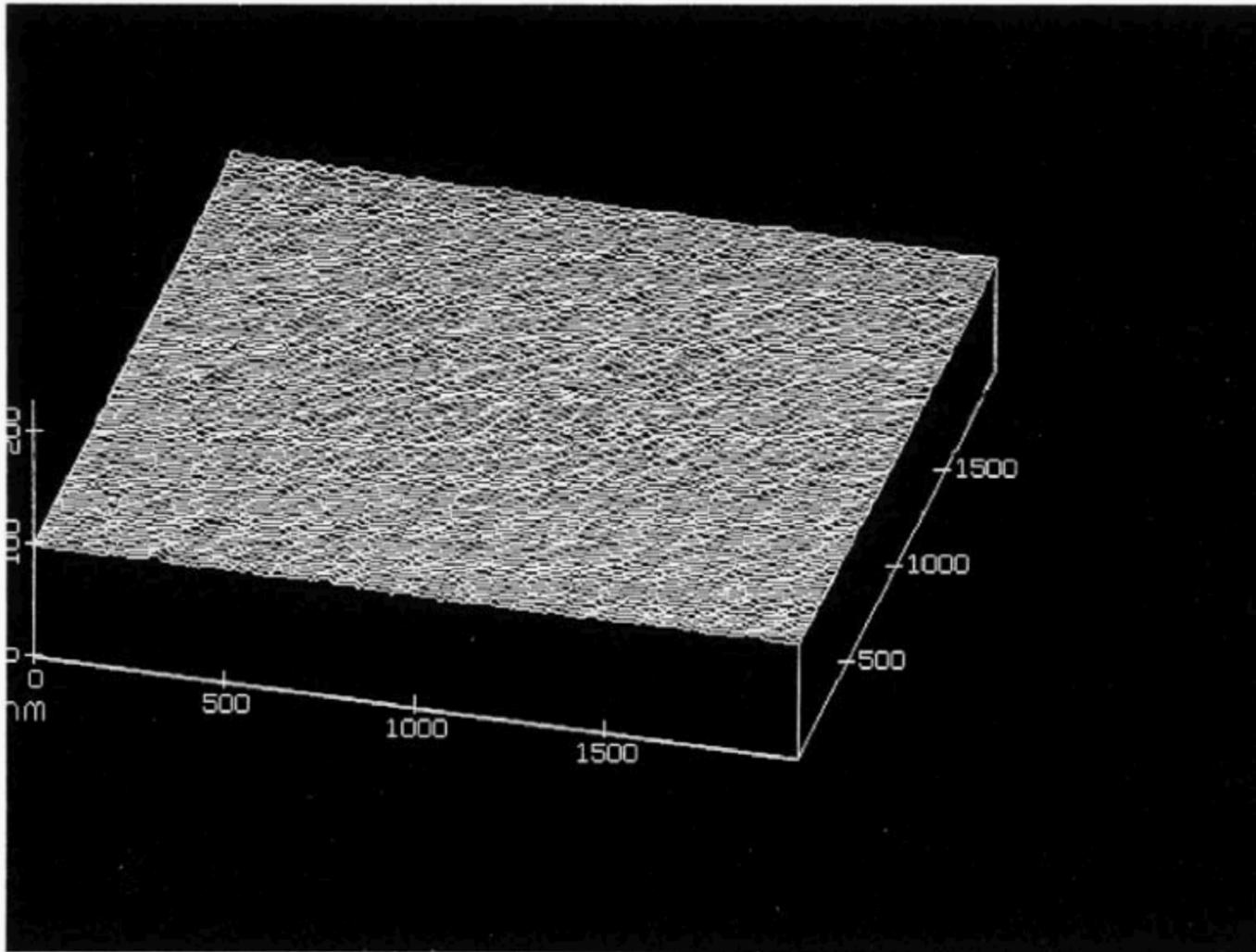


FIG. 6 Line Plot of the Same Scan as Shown in Fig. 5

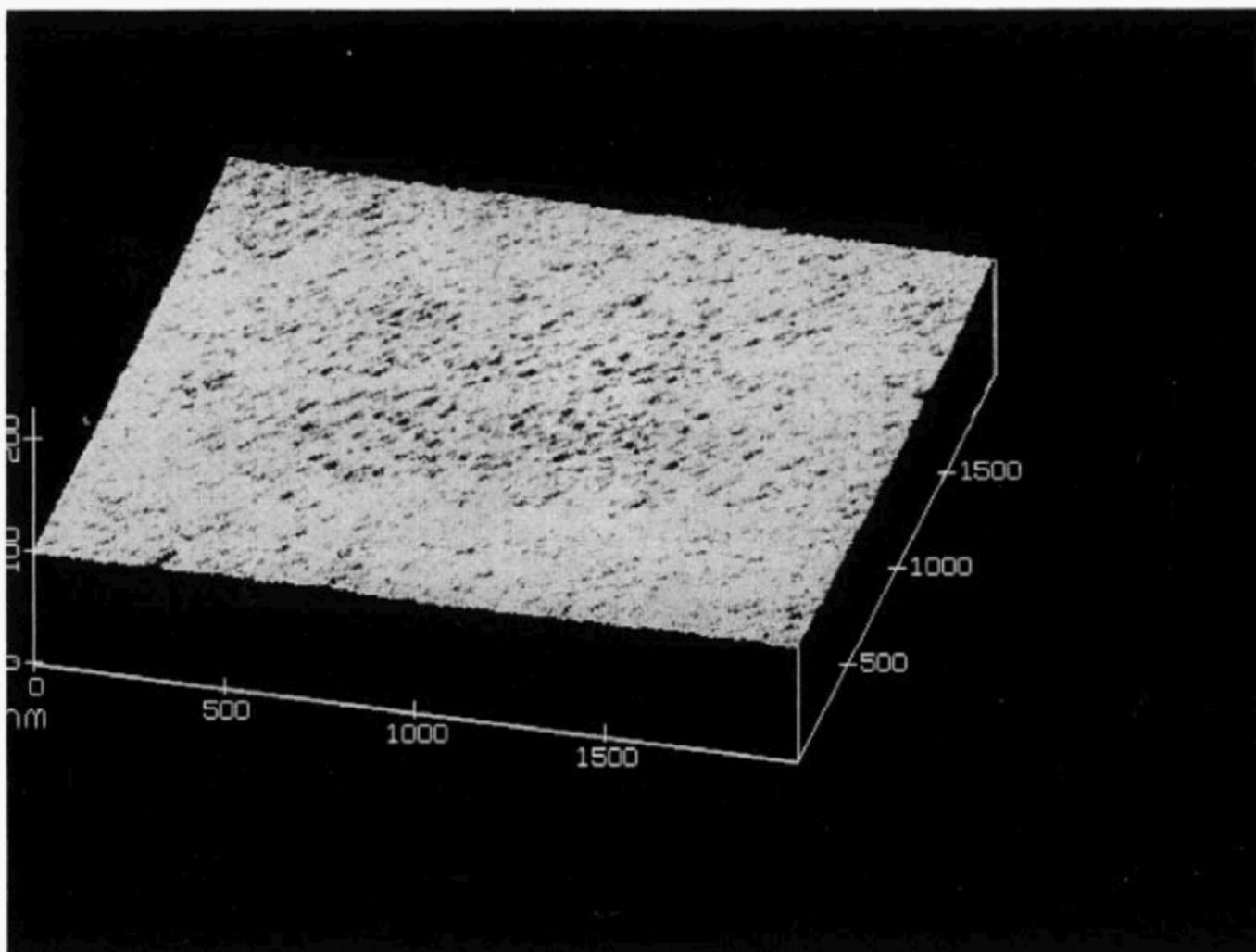


FIG. 7 Top Illuminated (Tilted Surface) View of Same Scan as Shown in Figs. 5 and 6

12.9 The following are the equations describing the parameters mentioned in 12.3. All surface data (Z values) are in nanometers.

12.10 *Surface Calculations:*

12.10.1

$$\text{Root - Mean - Square} = Z_{\text{rms}} = \left[\left(\frac{1}{N} \right) \sum_{i=1}^N (Z_i - \bar{Z})^2 \right]^{1/2}$$

where:

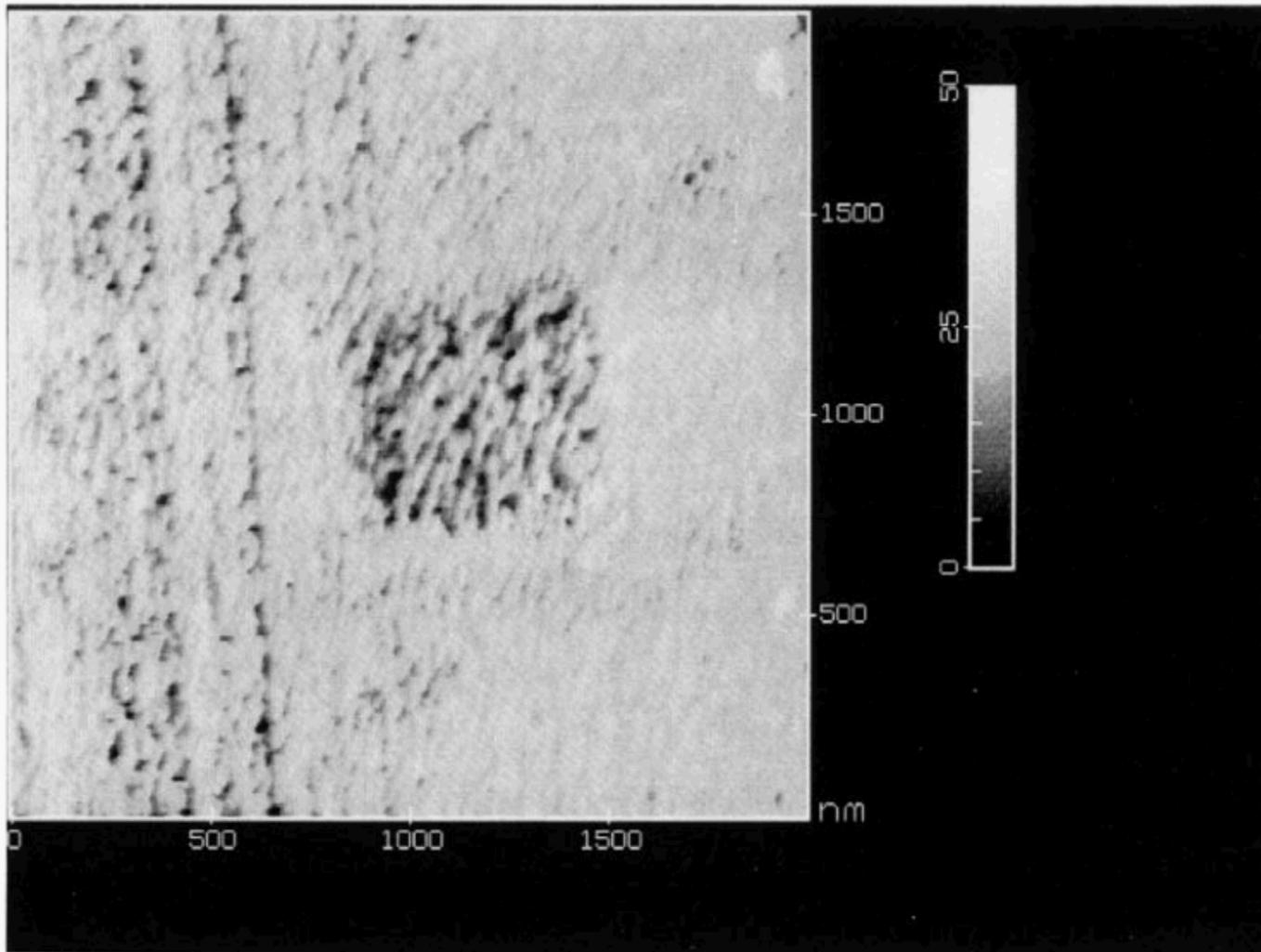


FIG. 8 Image of 2000-nm (Shaded Height Topview) Showing Damaged Area from Preceding 500-nm Scan. This Surface is Not Conductive Enough to Allow Surface Analysis by This Method.

TABLE 1 Summary of Figures of Merit

Identification	R_a (nm)	R_z (nm)	R_{max} (nm)	SA Index ^A	Z_{max} (nm)	Z_{rms} (nm)
Fig. 1, 500 nm scan	0.71 ± 0.14	3.1 ± 0.68	4.7 ± 1.6	20	9.1	1.0
Fig. 5, 2000 nm scan	0.74 ± 0.10	4.6 ± 0.92	5.6 ± 1.3	18	16	1.9
Fig. 9, ^B (a) 20 μ m scan	190	3562	630
Fig. 9, ^B (b) 20 μ m scan	11	470	500
Fig. 9, ^B (c) 20 μ m scan	0.19	190	40

^A See Appendix X1.

^B Profile data unavailable.

$$\bar{Z} = \left(\frac{1}{N} \sum_{i=1}^N Z_i \right)$$

where:

Z_i = height at point (x, y),

i = number of measurement, and

N = number of data points over area.

12.10.2

$$Z_{max} = Z_{largest} - Z_{smallest}$$

where:

$Z_{largest}$ = Z_i with the largest value (nm) over an area, and

$Z_{smallest}$ = Z_i with the smallest value (nm) over an area.

12.11 Profile Calculations:

12.11.1

$$R_a \text{ (mean roughness)} = \left(\frac{1}{N} \sum_{i=1}^N |Z_i - \bar{Z}| \right)$$

where:

Z_i = height at point (x, y) over a single profile,

i = number of the measurement in the profile, and

N = number of data points over entire profile.

12.11.2 R_{max} = largest peak-to-valley difference over profile length.

12.11.3 R_z = 10 point height

$$R_z = \frac{1}{5} \left(\sum_{i=1}^5 P_i + \sum_{i=1}^5 V_i \right)$$

where:

P_i = i th highest peak over the entire profile length, and

V_i = i th deepest valley over the entire profile length.

13. Interpretation of Data

13.1 This test method only allows digital filtration of the data for purposes of numerical analysis and data presentation.

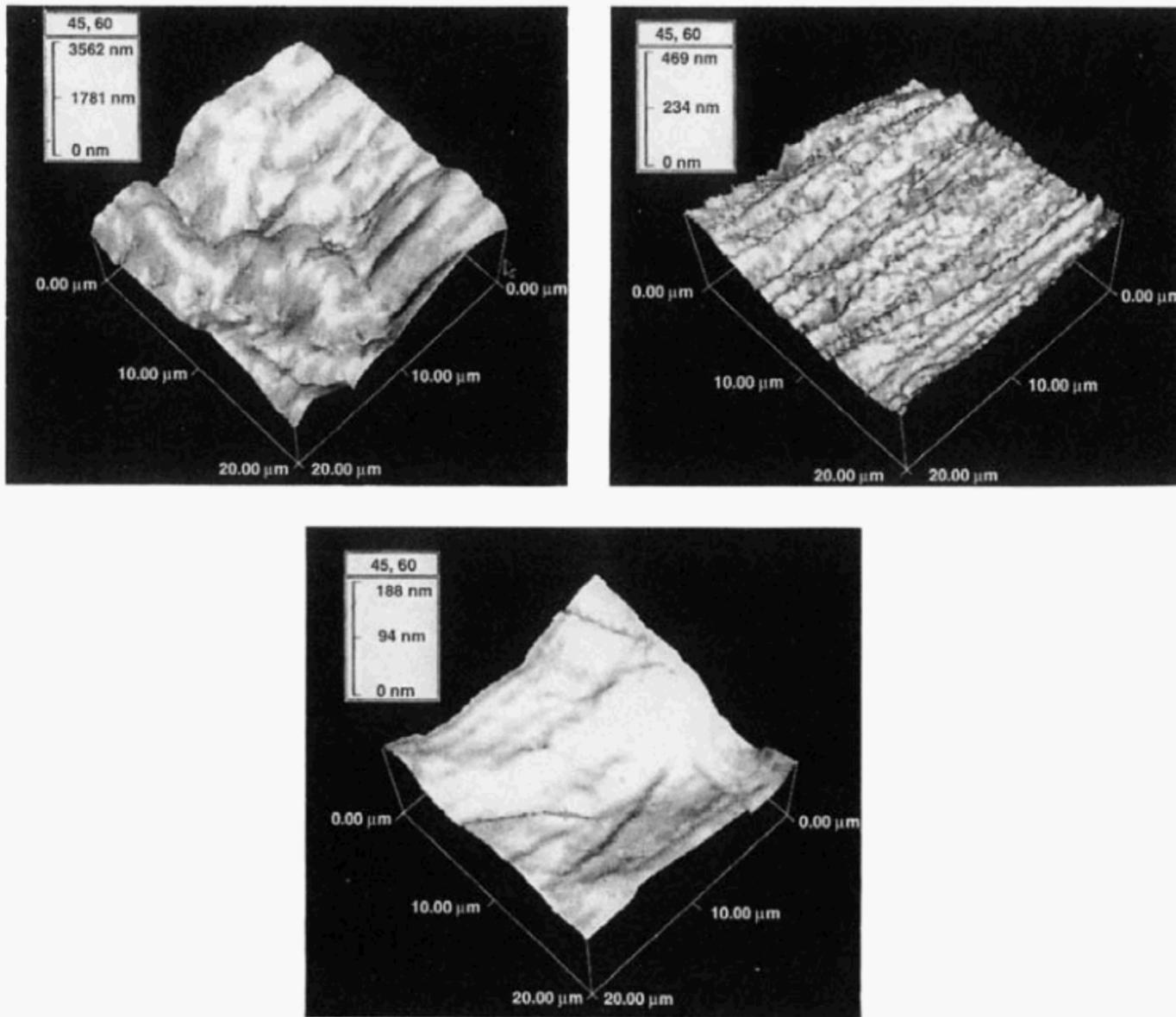


FIG. 9 Stainless Steel Surfaces Used for SA Index, Z_{max} , and Z_{rms} Data Given in Table 1

13.1.1 Correction for scanning arc by plane-fitting or high frequency noise by low-pass filtering are allowed only for image appearance.

13.1.2 Frequency filtering prior to calculation of figures of merit is only allowed if the definition of that figure requires it (see Calculation Section). Surface figures of merit can be determined from plane-fitted data.

13.1.3 Present all images corresponding to data used for figures of merit.

13.2 The data for all scan area sizes will be shown as surface topography by plotting the feature height (Z_i) as a function of the tip location in the x - y plane.

13.3 The method of representing the surface will be either (1) a shaded height topview, or (2) a line plot. If the instrument's imaging capabilities cannot provide a shaded height plot, then the line plot can be given.

13.4 If both representations are possible, the shaded height plot will be the primary choice.

13.5 The full height scale value for the shaded plot will be 50 nm regardless of scan width. Fig. 1 shows an example of a 500-nm wide area with heights plotted on a scale from 0 to 50 nm. Fig. 5 shows the same representation for a 2000-nm wide area.

13.6 Other image representations can also be used, in addition to the shaded height or line plots, for the sake of clarity or to overcome photocopying difficulties.

13.7 The tilted surface line plot will be plotted as shown in Fig. 2 (500 nm) and 6 (2000 nm). The angle of tilt and rotation can be arbitrary. The height scale (z -direction) will be exaggerated (magnified) with respect to the scan width (x -direction) by a factor of $3 \pm 10\%$, such that:

$$\frac{\text{nanometers of scan width per centimeter as printed}}{\text{nanometers of height scale per centimeter as printed}} = 3 \pm 0.3$$

13.8 Changes in scan width will be factored into the height scale for images represented in a manner other than a shaded topview (for example, illuminated surface or line plot). Fig. 5 shows a 2000-nm wide area (four times the width of Fig. 1) with heights plotted on a scale from 0–200 nm (four times the height scale of Fig. 1).

13.9 Surface profiles should also conserve this scaling between scan width.

13.10 Also included for clarity are Fig. 3 and Fig. 7, which are illuminated tilted surfaces for the same two areas already shown.

13.11 Fig. 9 shows different surfaces obtained from another instrument, also using the illuminated surface representation.

14. Precision and Bias

14.1 The primary determination of this test method is graphical and not numerical, hence no statement is made about the precision or bias of these results.

14.2 *Precision*—The precision of the secondary, (suggested) numerical results (for both profiles and areas) as determined by the statistical examination of interlaboratory test results are as follows:

14.2.1 *Repeatability*— The repeatability standard deviation (RSD) for successive results obtained by the same operator for the same sample using the same apparatus, in the normal and correct operation of the method is as follows:

	RSD, %
Z _{max}	39
R _a	31

14.2.2 *Reproducibility*— to be determined.

NOTE 2—These values were determined in a program in compliance with Practice E691.

14.3 *Bias*—Bias depends on the empirical conditions of this test method.

15. Keywords

15.1 average roughness; clean room; components; microscope; profile; roughness; scanning; stainless steel; surface; texture; topography; tunneling

APPENDIXES

(Nonmandatory Information)

X1. SUGGESTED OPERATING CONDITIONS FOR ELECTROPOLISHED STAINLESS STEEL SURFACES

X1.1 The images in Figs. 1 through 8 were obtained for 0.25-in. stainless steel tubing, with tip voltage at 1800 mV, current levels monitored at 0.5 nA in constant current mode, and the scan rate set at about 2 s/μm in the x-direction. The tip was Pt/Ir. The tubing was cut, first longitudinally and then transversely, to a length of 0.5 cm, and briefly cleaned with

ethyl alcohol. The sample was imaged in air. The longitudinal direction of the tubing sample is in the y-direction in the image. A plane fitting filter was applied to each image. In some cases, the contrast of the images was increased for photoduplication purposes.

X2. SURFACE AREA TERMINOLOGY

X2.1 Definitions:

X2.1.1 *ideal surface area*—area of least-squares fitted plane for the surface analyzed.

X2.1.2 *SA index*—see **surface area index**.

X2.1.3 *surface area index (SA index)* —the area of a best fit plane (or ideal surface) subtracted from the actual area calculated for the surface, divided by the ideal area, and multiplied by 1000 (see Surface Calculations in X2.2.1).

X2.1.4 *true surface area*—actual surface area, being greater than the ideal surface area due to surface features such as peaks and valleys.

X2.1.5 *Surface Calculations*:

X2.1.6 *Surface Area Index*:

$$\text{SA Index} = \frac{(\text{True surface area} - \text{Ideal surface area})}{\text{Ideal Surface Area}} \times 1000$$

Ideal surface area = area of least-squares fitted plane

ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.

This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.

This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org). Permission rights to photocopy the standard may also be secured from the Copyright Clearance Center, 222 Rosewood Drive, Danvers, MA 01923, Tel: (978) 646-2600; http://www.copyright.com/