



Chain of Custody: 300396
Client: US Food & Drug Administration
Address: Office of Cosmetics & Colors
 4300 River Road
 College Park, MD 20740
Attention: John Gasper

Job Name: Task 3 - Analysis of Official Samples
Job Location: 1st Group - 8 Samples
Job Number: CLIN 1 - Task 3 (8 Samples)
PO Number: HHSF223201810337P

Date Submitted: 3/14/2019
Date Analyzed: 3/29/2019 - 4/18/2019
Report Date: 4/25/2019
Date Sampled: Not Provided
Person Submitting: Steve Wolfgang
Revised: 4/30/2019 (1st Revision)

SUMMARY OF ASBESTOS IN TALC ANALYSIS

AMA Sample ID	Client Sample ID	TEM LOD Using ASTM D5756 Mass Calculation	TEM LOQ Using ASTM D5756 Mass Calculation	% Tremolite by TEM Using ASTM D5756 Mass Calculation	% Chrysotile by TEM Using ASTM D5756 Mass Calculation	% Total Asbestos a by TEM Using ASTM D5756 Mass Calculation	Asbestos by PLM	% Organics	% Acid Soluble	% Other	Comments
300396-1	D-32	0.00000218%	0.00000872%	ND	ND	ND	ND	6.2%	5.0%	88.8%	
300396-1A	D-32	0.00000162%	0.00000648%	ND	ND	ND	ND	6.2%	1.4%	92.4%	
300396-1B	D-32	0.00000144%	0.00000574%	ND	ND	ND	ND	6.2%	4.0%	89.8%	
300396-2	D-33	0.00000192%	0.00000769%	ND	ND	ND	ND	28.7%	3.3%	68.0%	
300396-2A	D-33	0.00000205%	0.00000819%	ND	ND	ND	ND	29.4%	2.1%	68.5%	
300396-2B	D-33	0.00000193%	0.00000773%	ND	ND	ND	ND	29.1%	3.3%	67.6%	
300396-3	D-34	0.00000254%	0.00001016%	ND	ND	ND	ND	24.7%	4.7%	70.6%	
300396-3A	D-34	0.00000285%	0.00000744%	ND	< 0.00080%	< 0.00080%	ND	23.4%	4.5%	72.1%	
300396-3B	D-34	0.00000370%	0.00001479%	ND	0.00030%	0.00030%	ND	24.0%	3.9%	72.0%	
300396-4	D-35	0.00000134%	0.00000536%	0.00071%	0.00503%	0.00574%	ND	12.8%	12.4%	74.7%	
300396-4A	D-35	0.00000188%	0.00012905%	< 0.00013%	< 0.00013%	< 0.00013%	ND	13.7%	13.8%	72.5%	
300396-4B	D-35	0.00000168%	0.00000671%	0.00367%	0.00005%	0.00371%	ND	12.5%	15.5%	73.0%	
300396-5	D-36	0.00000188%	0.00000751%	ND	ND	ND	ND	24.0%	3.5%	72.5%	
300396-5A	D-36	0.00000114%	0.00000454%	ND	ND	ND	ND	24.1%	3.0%	72.8%	
300396-5B	D-36	0.00000150%	0.00000599%	ND	ND	ND	ND	24.1%	2.5%	73.4%	
300396-6	D-37	0.00000150%	0.00000599%	ND	ND	ND	ND	19.2%	6.7%	74.1%	
300396-6A	D-37	0.00000178%	0.00000714%	ND	ND	ND	ND	18.9%	5.3%	75.8%	
300396-6B	D-37	0.00000157%	0.00000629%	ND	ND	ND	ND	18.3%	6.0%	75.8%	
300396-7	D-38	0.00000134%	0.00000536%	ND	ND	ND	ND	0.0%	3.1%	96.8%	
300396-7A	D-38	0.00000173%	0.00000694%	ND	ND	ND	ND	0.1%	2.4%	97.5%	
300396-7B	D-38	0.00000135%	0.00000539%	ND	ND	ND	ND	2.5%	2.5%	97.4%	
300396-8	D-39	0.00000131%	0.00000524%	ND	ND	ND	ND	55.3%	11.0%	33.7%	
300396-8A	D-39	0.00000180%	0.00000721%	ND	ND	ND	ND	55.3%	8.3%	36.4%	
300396-8B	D-39	0.00000135%	0.00000540%	ND	ND	ND	ND	55.3%	15.4%	29.3%	

LOD = Limit of Detection **LOQ** = Limit of Quantification **ND** = Not Detected **PLM** = Polarized Light Microscopy **TEM** = Transmission Electron Microscopy

Analytical Method(s): PLM by Modified NY ELAP 198.6
 TEM by Modified NY ELAP 198.4/ASTM D5756

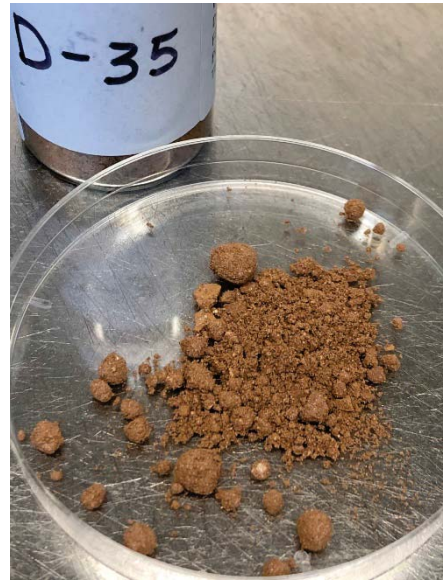
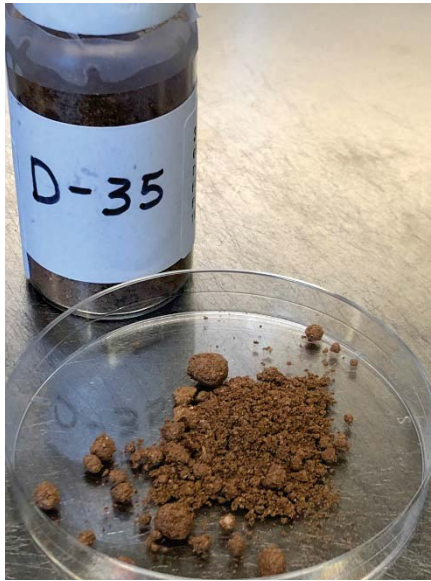
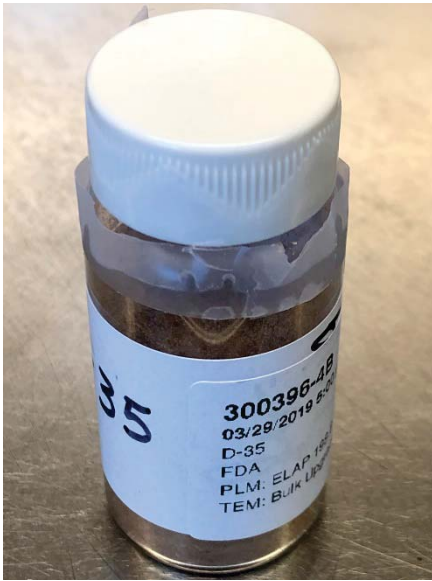
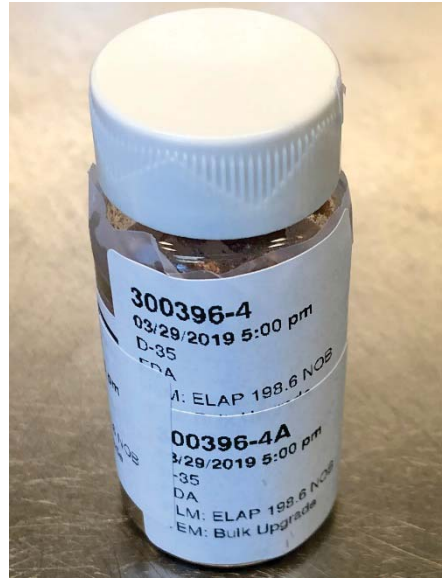
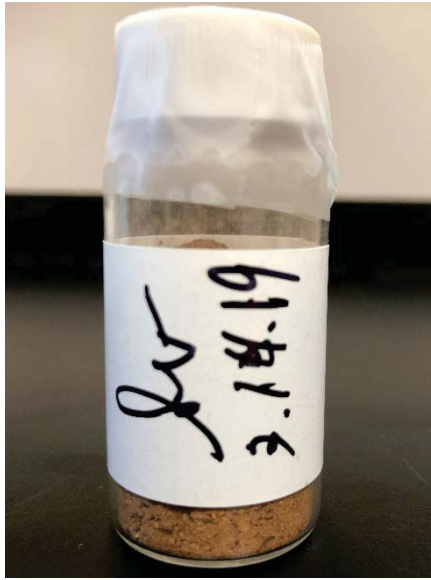
Analyst(s): PLM
 TEM

[Signature]
Technical Director: Andreas Saldivar

All results are to be considered preliminary and subject to change unless signed by the Technical Director or Deputy

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300396-4, 4A, 4B/D-35



(b) (4)

(b) (4)

Sample Preparation

Samples were prepared for PLM and TEM bulk analysis by (b) (6) on March 15, 2019 through March 29, 2019.

Sample preparation consisted of the following steps:

- 1) Label and weigh two 8mL glass vials for each sample in the set – one vial for the PLM preparation and one vial for the TEM preparation.
- 2) Weigh out 0.1 to 0.8 grams of material and place in corresponding 8mL glass vial. Record weight.
- 3) Burn samples at 480° C for at least 12 hours.
- 4) Record Post-Ash Weight.
- 5) Treat ashed sample with concentrated hydrochloric acid.
- 6) Filter acid reduced material onto a pre-weighed 47mm 0.4um PolyCarbonate filter.
- 7) Place filter into drying oven for 30 minutes and then record Post-Acid Reduced weight.
- 8) Make four PLM slide preparations from the PLM residual ash for each sample in 1.550 dispersion oil. Make additional preparations in 1.605, 1.625, 1.680 and 1.700 dispersion oil as necessary for particle identification.



- 9) Weigh a portion of the residue from the TEM residual ash and place it into the corresponding pre-weighed 100ml jar.
- 10) Fill the 100ml jar with deionized water
- 11) Sonicate the jars for approximate 5-minutes.
- 12) Filter 0.2ml to 1ml of the solution onto a 47mm 0.22um MCE filter.
- 13) Dry the filter for 10 minutes then collapse, carbon coat, and place on a 3 TEM grids.

PLM Analysis

Analysis was performed in accordance with NY ELAP 198.6 protocols. The analysis was conducted using an Olympus BH-2 polarized light microscope (PLM) equipped with a dispersion staining objective. All four slide preparations for each aliquot were examined. 400-point count was performed for those samples on which asbestos was observed. If no asbestos was detected on any of the slides, the percentage of fibrous components was determined by visual estimation. The results of this analysis are detailed below in the *Discussion and Interpretation of Analytical Findings* section for each individual sample.

TEM Analysis

Analysis was performed in accordance with modified NY ELAP Method 198.4 protocols. The analysis was performed using a JEOL JEM-100CX II transmission electron microscope (TEM), equipped with a Thermo Fisher Quest Energy Dispersive X-Ray Analyzer (EDXA), at magnifications of 19,000x. Two grids for each aliquot were examined. Twenty (20) grid openings were examined per sample.

Modifications to the NY ELAP 198.4 Method were:

- 1) The residue was not placed in alcohol and prepared using the quick drop method. To obtain a more uniform preparation, the residue was placed in a jar and filled with 100ml of deionized water. The jar was sonicated, and a portion of the solution was filtered onto a 47mm 0.22um MCE filter.
- 2) The tremolite and chrysotile were not visually estimated. The length and width of the observed particles were measured, and the mass of each amphibole particle was calculated using the ASTM D5756 method.

The results of this analysis are detailed below in the *Discussion and Interpretation of Analytical Findings* section for each individual sample.

Calculations

ASTM D5756 Mass

$$M = \pi/4 L * W^2 * D * 10^{-12}$$

M = mass

L = length

W = width

D = density

Percent Calculation

$$\frac{EFA(\text{mm}^2) * 100\text{ml} * MA(\text{g}) * RW(\text{g})}{VF(\text{ml}) * IW(\text{g}) * AA(\text{mm}^2) * RJ(\text{g})}$$

The calculated value is then multiplied by 100 to convert it to percent.

EFA – Effective filter area

MA – Mass of asbestos

RW – Weight of residue

VF – Volume filtered

IW – Initial weight of the sample

AA – Area analyzed

RJ – Weight of residue placed into the jar



Limit of Detection and Quantification

We used the mass of a 0.5 x 0.04-micron tremolite or chrysotile fiber, depending on what was found in each sample, as the basis for our calculations. Limit of detection was defined as 1 fiber and limit of quantification was defined as 4 fibers.

Some aliquots of samples (b) (4) D35 contained very small amounts of asbestos that were either at or below our 4-fiber limit of quantification. For these samples we defined our limit of quantification as follows:

300396-3A: mass of the single observed chrysotile fiber plus the mass of three tremolite fibers measuring 0.5 x 0.04 microns

300396-4A: mass of the two observed chrysotile fibers, the single observed tremolite structure plus the mass of one 0.5 x 0.04 microns tremolite fiber.

Discussion and Interpretation of Analytical Findings:

(b) (4) [Redacted]

[Redacted]

[Redacted]

[Redacted]

(b) (4) [Redacted]

(b) (4)

(b) (4)

300396-4, 4A, 4B, Client Sample D35

PLM
All three aliquots of sample D35 were analyzed by (b) (6) on March 29, 2019. No asbestos or non-asbestos amphibole variants were detected the samples. The results were calculated using the equations detailed in the calculations section.

300396-4	NAD
300396-4A	NAD
300396-4B	NAD

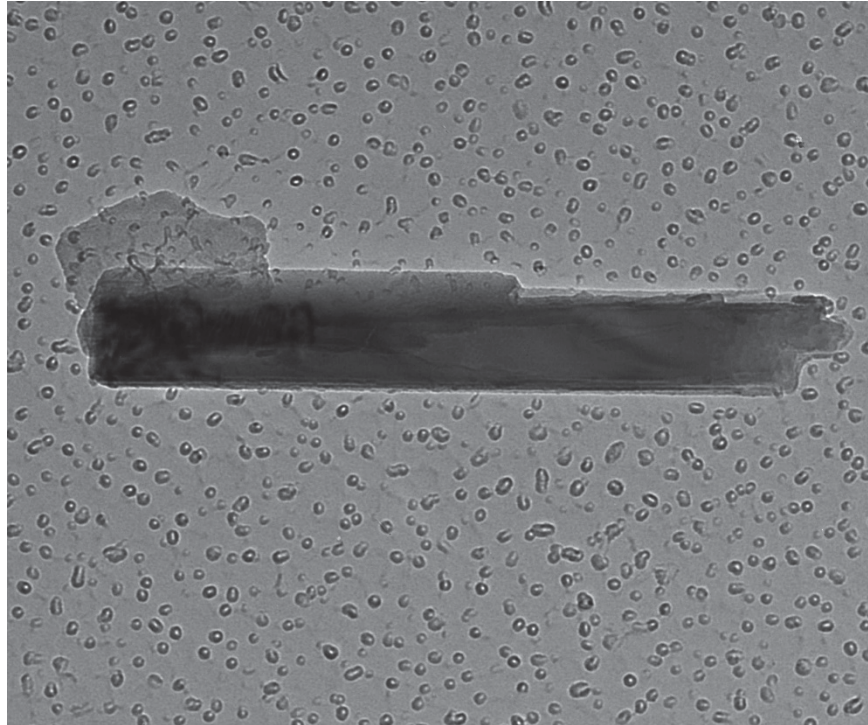
TEM
Samples 4 and 4A were analyzed by (b) (6) on April 11, 2019. He analyzed sample 4B on April 16, 2019. The primary particles observed were mica and talc. Some titanium particles were observed. The talc was mostly flakes but there were also some ribbons and fibers. Tremolite and chrysotile were observed and counted on all three samples. Two tremolite and two chrysotile structures were counted on sample 4. One tremolite and two chrysotile structures were counted on sample 4A. One tremolite and five chrysotile structures were counted on sample 4B. The results were calculated using the equations detailed in the calculations section.

300396-4	0.00574%
300396-4A	< 0.00013%
300396-4B	0.00371%

Below are pictures, diffraction patterns, and chemistry of the counted tremolite and chrysotile particles. Some of the talc fibers and ribbons are also pictured below. The unidentified peaks in chemistry spectra are copper, zinc, and carbon. Those peaks are from the TEM specimen holder and specimen grid.



Sample 300396-4 Tremolite fiber #1

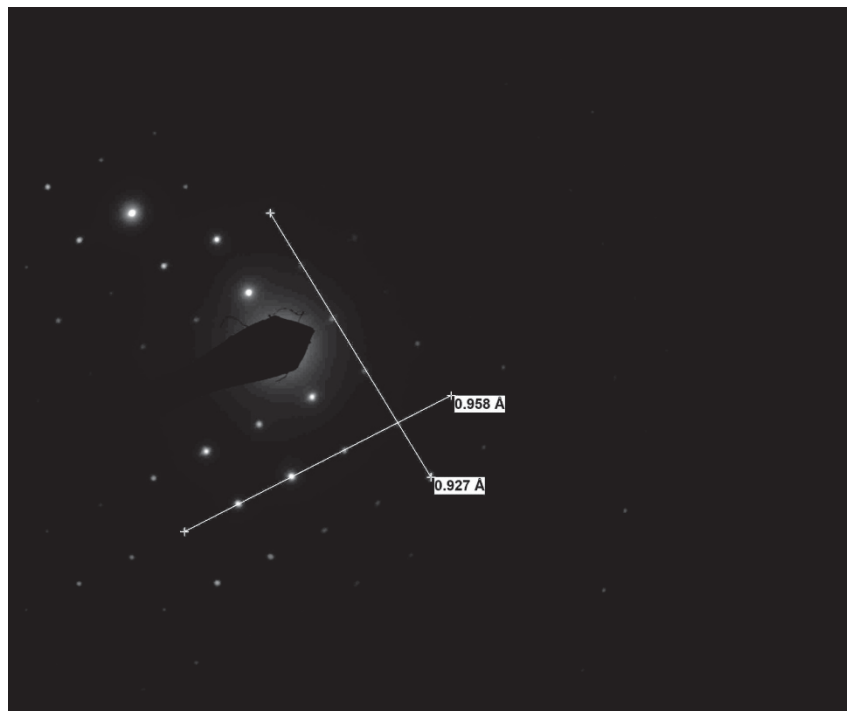


300396 FDA_072.jpg
Tremolite Fiber
Cal: 0.541520 nm/pix
10:44 4/11/2019

100 nm
HV=100kV
Direct Mag: 19000 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4 Zone axis diffraction pattern from tremolite fiber #1 pictured above.

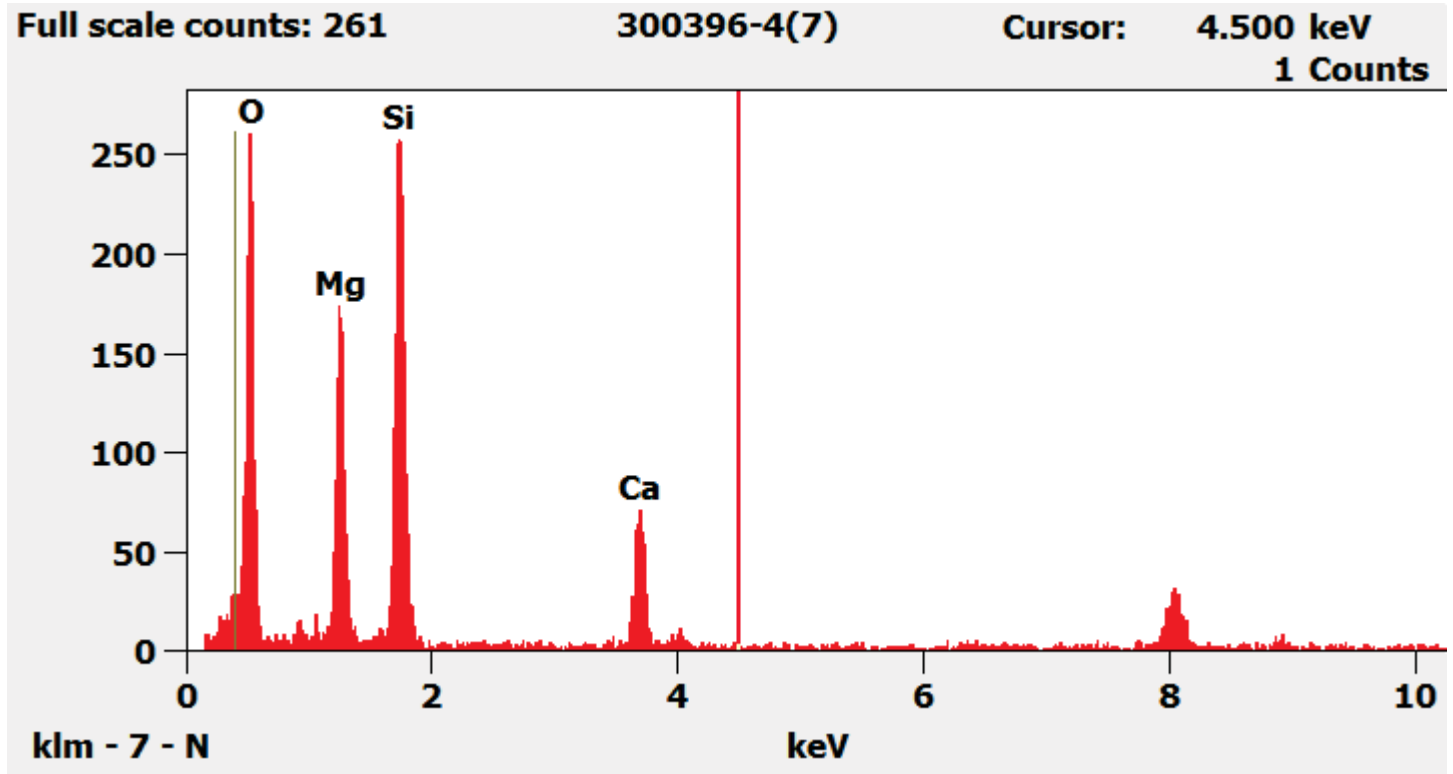


300396 FDA_074.jpg
Tremolite Fiber Zone Axis
[0 2 -2]
10:49 4/11/2019

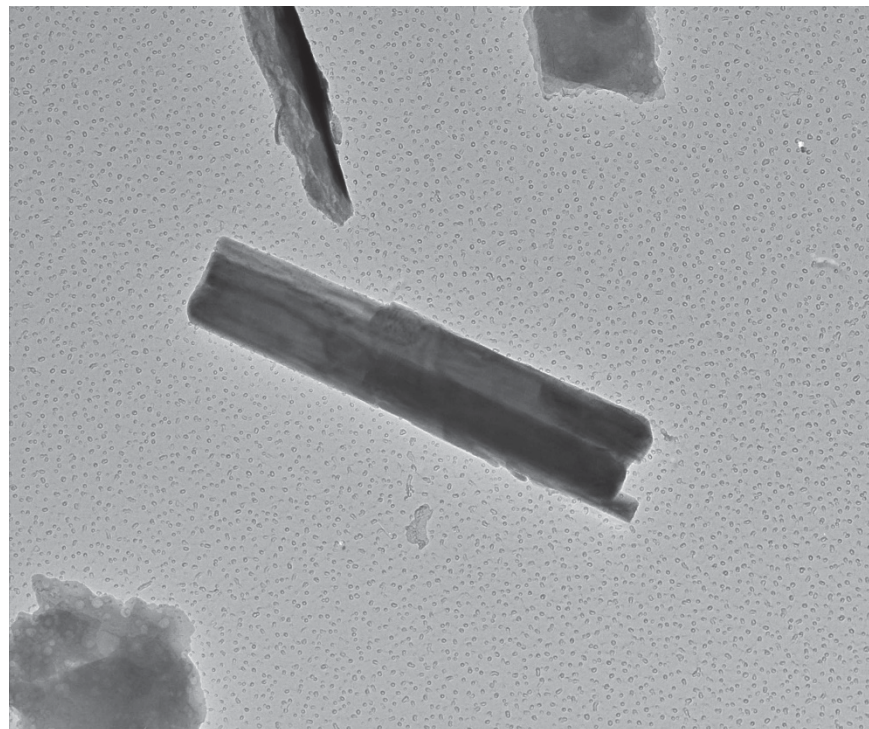
100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4 Chemistry from tremolite fiber #1



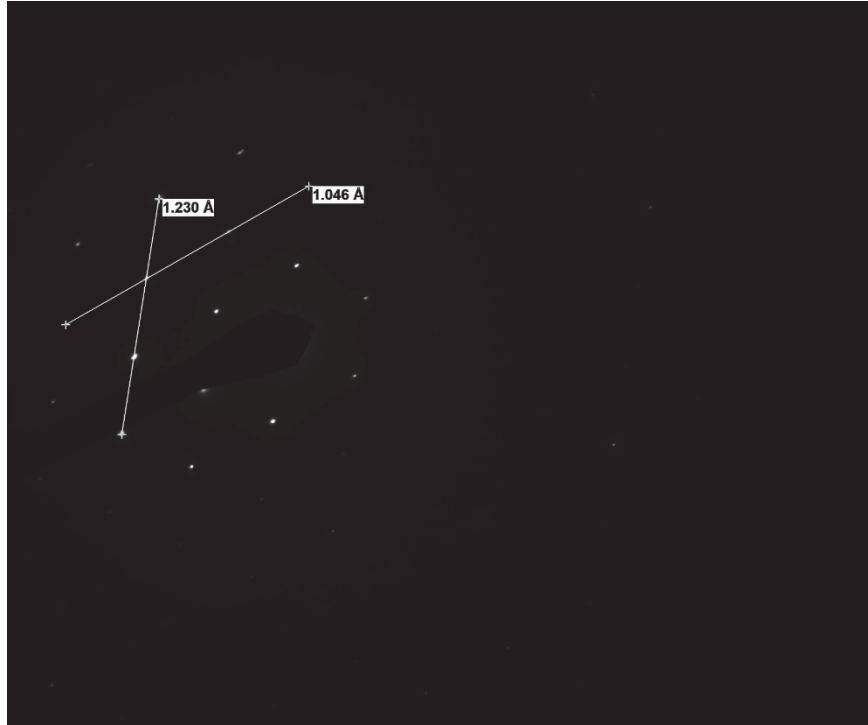
Sample 300396-4 Tremolite fiber 2



300396 FDA_089.jpg
Tremolite Fiber
[-1 1 -4]
Cal: 0.001429 µm/pix
12:34 4/11/2019
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

400 nm
HV=100kV
Direct Mag: 7200 x
AMA Analytical Services, Inc

Sample 300396-4 Diffraction patterns from the tremolite fiber #2 pictured above.



300396 FDA_088.jpg
Tremolite Zone Axis
[-1 1 -4]
12:28 4/11/2019

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

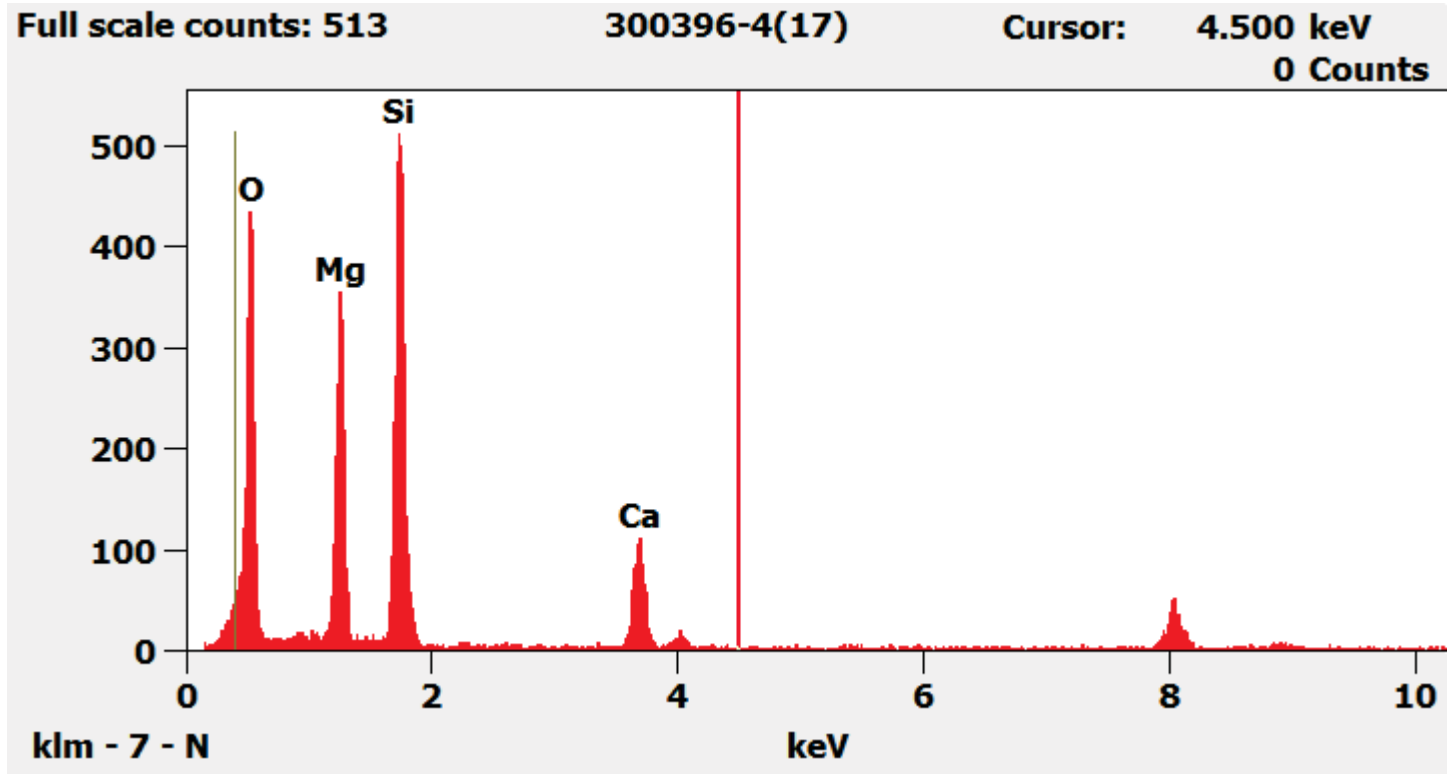


300396 FDA_087.jpg
Tremolite Diff
12:23 4/11/2019

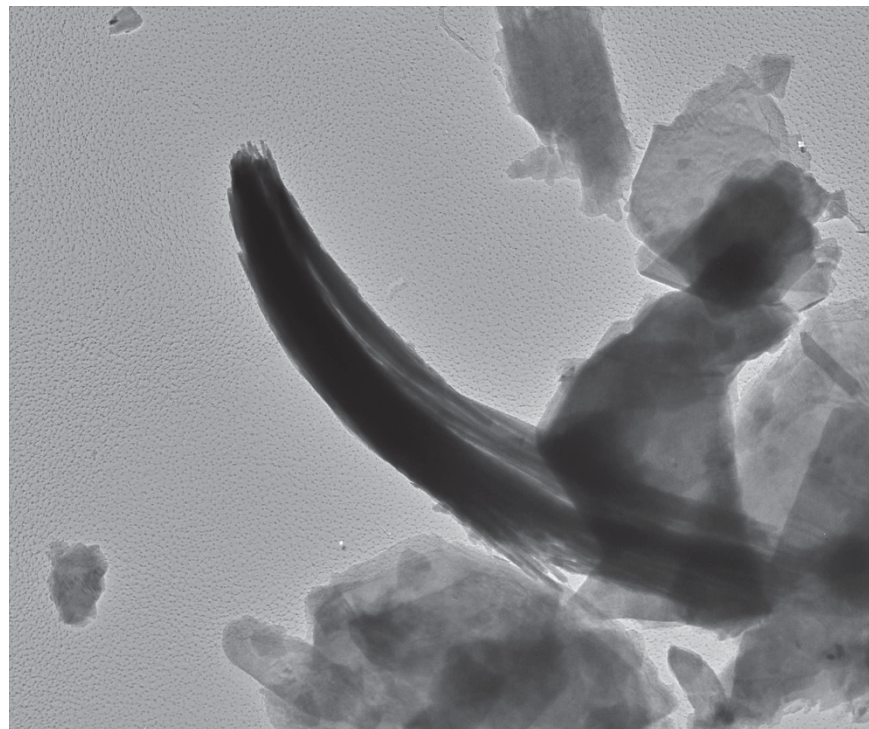
100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4 Chemistry from tremolite fiber pictured above.



Sample 300396-4 Chrysotile structure #1

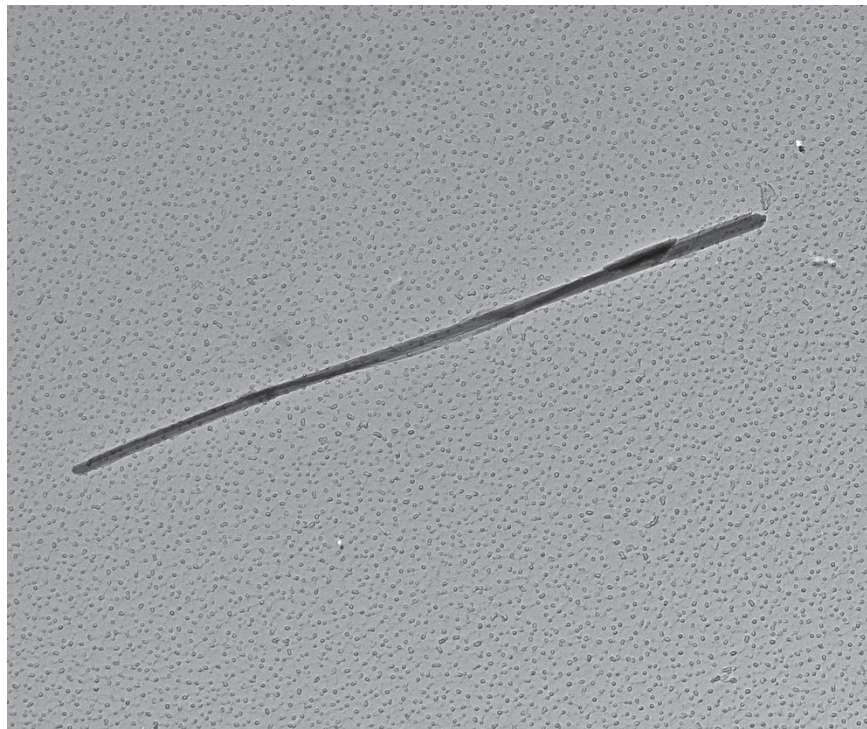


300396 FDA_075.jpg
Chrysotile Bundle
Cal: 0.002858 µm/pix
11:01 4/11/2019

800 nm
HV=100kV
Direct Mag: 3600 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4 Chrysotile structure #2



300396 FDA_083.jpg
Chrysotile Fiber
Cal: 0.001429 $\mu\text{m}/\text{pix}$
11:57 4/11/2019

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

400 nm
HV=100kV
Direct Mag: 7200 x
AMA Analytical Services, Inc

Sample 300396-4 Diffraction pattern from chrysotile structure #2 pictured above

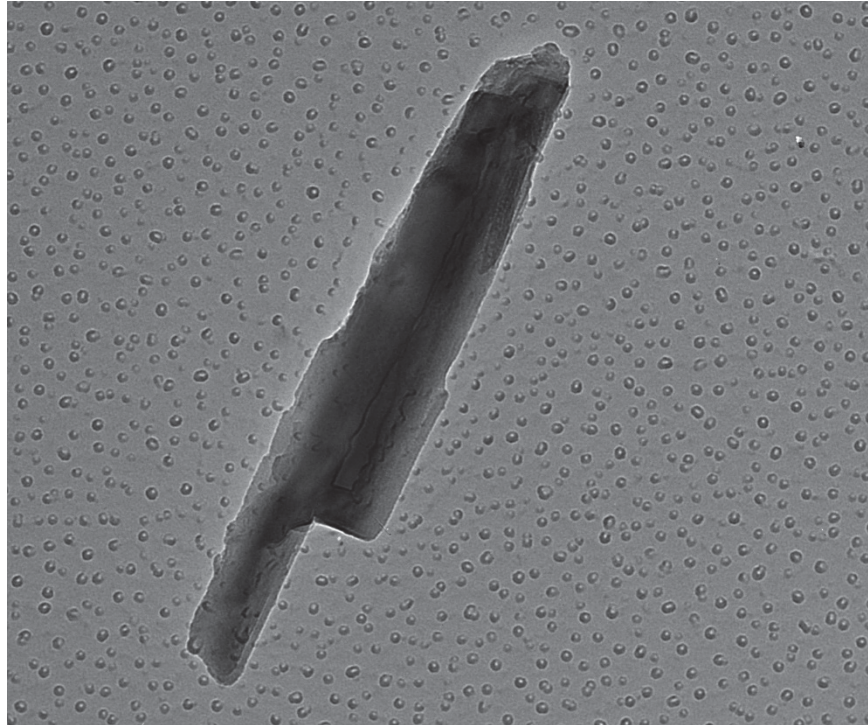


300396 FDA_082.jpg
Chrysotile Diffraction 2
11:46 4/11/2019

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

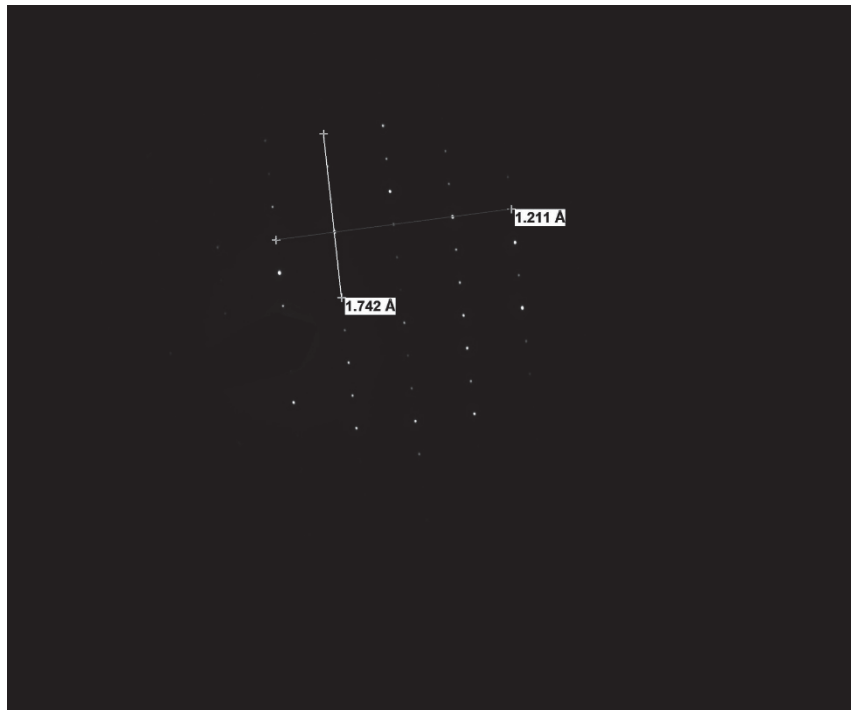
Sample 300396-4A Tremolite fiber



300396 FDA_096.jpg
Tremolite
[0 0 -4]
Cal: 0.541520 nm/pix
15:53 4/11/2019
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 nm
HV=100kV
Direct Mag: 19000 x
AMA Analytical Services, Inc

Sample 300396-4A Diffraction pattern from tremolite fiber pictured above.

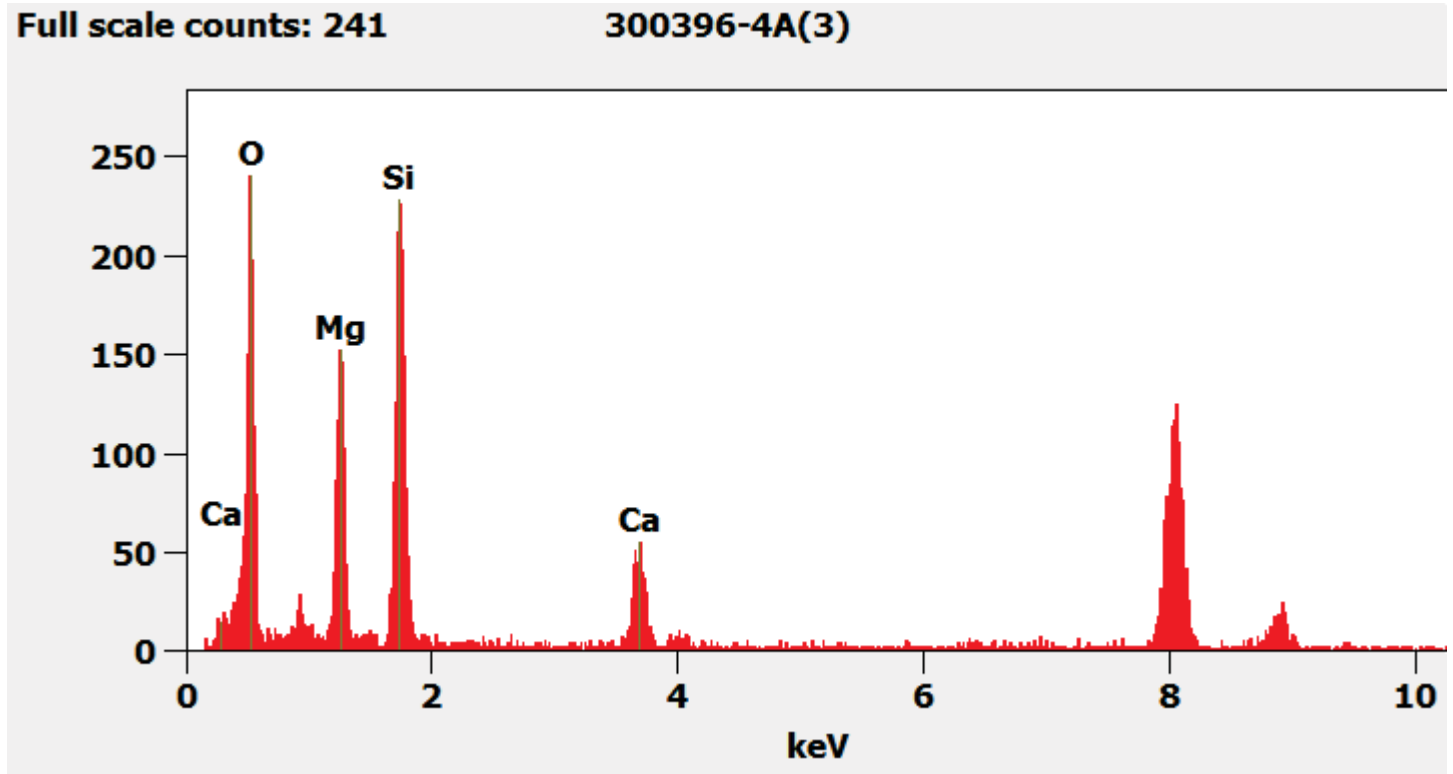


300396 FDA_095.jpg
Tremolite Zone Axis
[0 0 -4]
15:44 4/11/2019

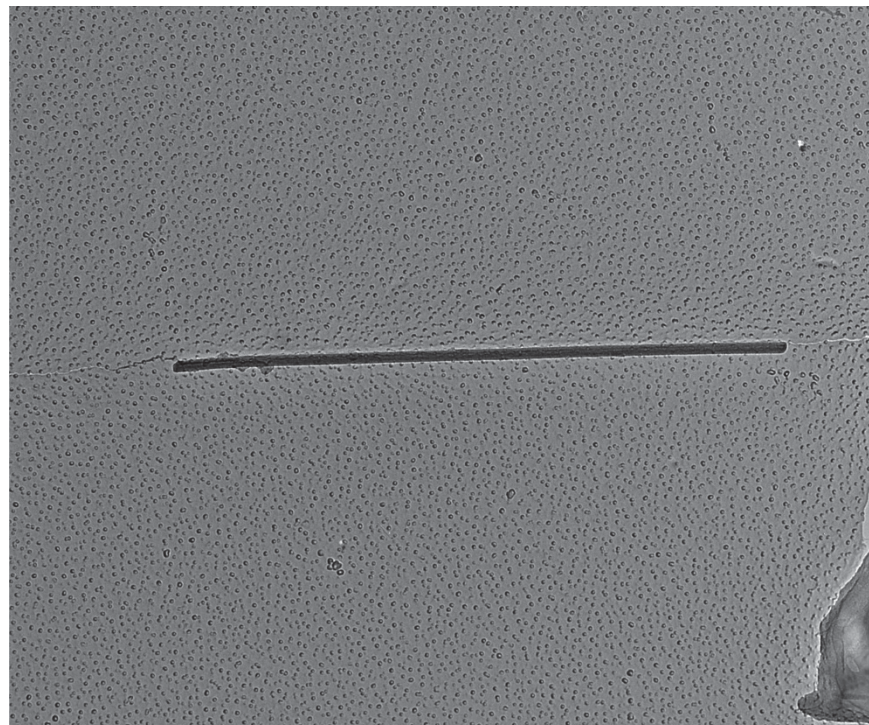
100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
Tilt:25
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4A Chemistry from tremolite fiber pictured above



Sample 300396-4A Chrysotile structure #1



300396 FDA_099.jpg
Chrysotile Fiber
Cal: 0.001429 $\mu\text{m}/\text{pix}$
16:10 4/11/2019

400 nm
HV=100kV
Direct Mag: 7200 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4A Diffraction pattern from chrysotile structure #1 pictured above

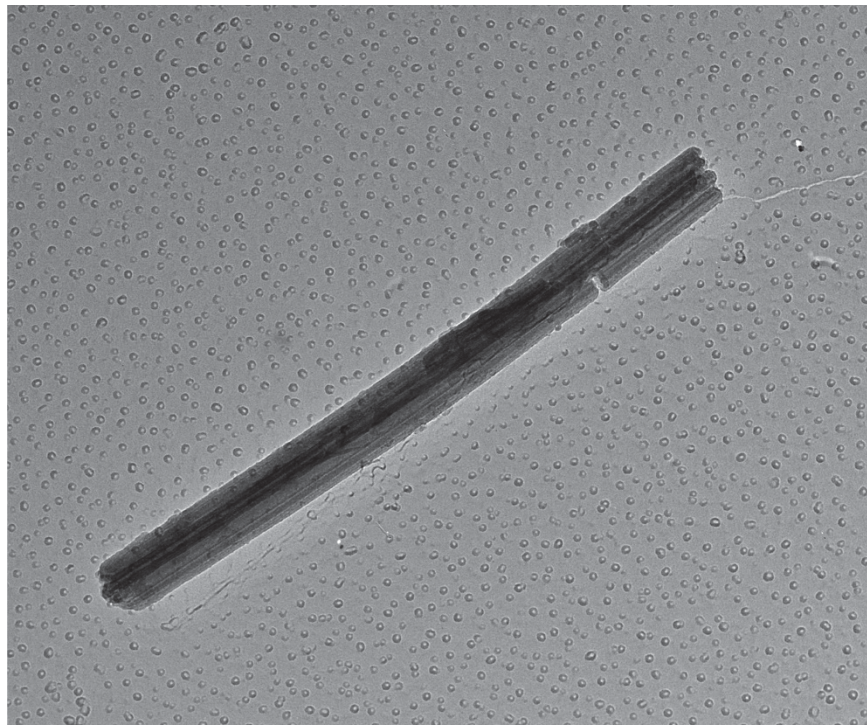


300396 FDA_097.jpg
Chrysotile
16:07 4/11/2019

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4A Chrysotile structure #2



300396 FDA_102.jpg
Chrysotile 2 Fiber
Cal: 0.734921 nm/pix
16:27 4/11/2019

200 nm
HV=100kV
Direct Mag: 14000 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4A Diffraction pattern from chrysotile structure #2 pictured above.

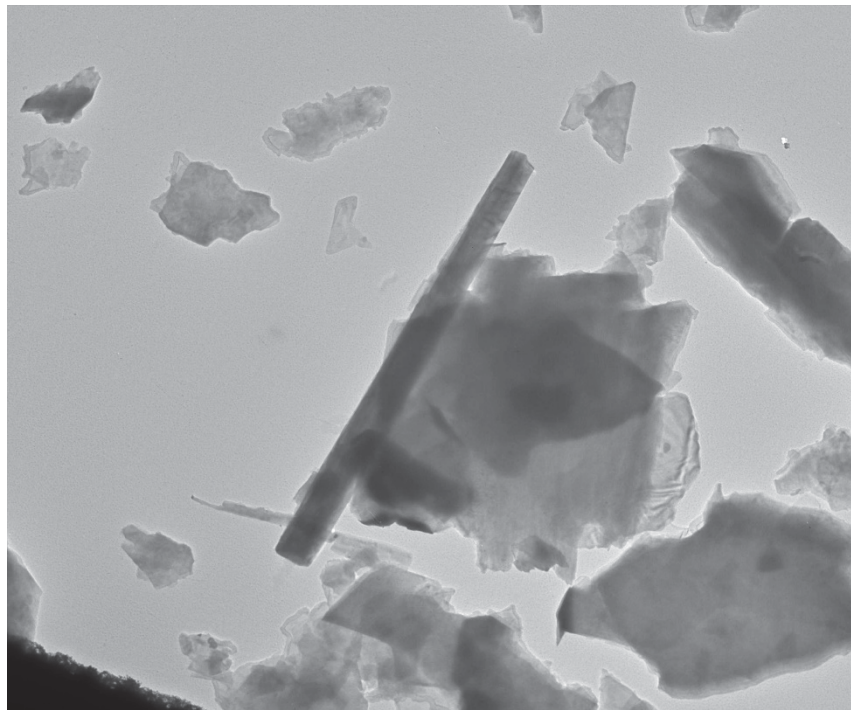


300396 FDA_100.jpg
Chrysotile 2 Diff
16:25 4/11/2019

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4B Tremolite fiber

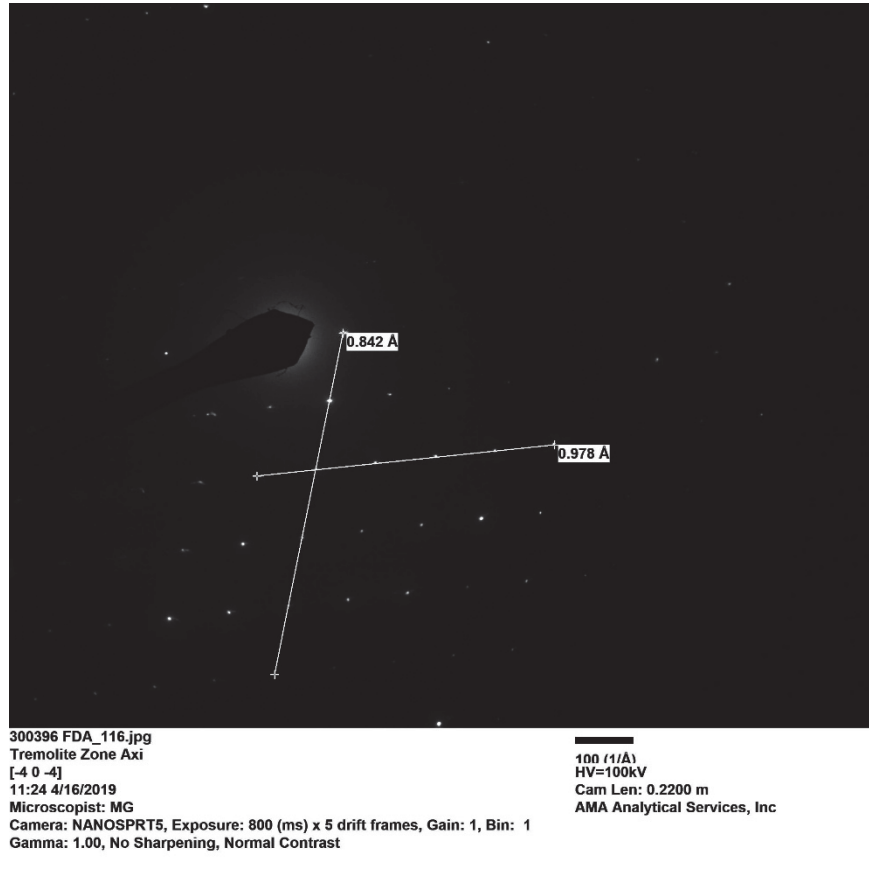


300396 FDA_117.jpg
Tremolite Fiber
[-4 0 -4]
Cal: 0.005415 µm/pix
11:30 4/16/2019
Microscopist: MG

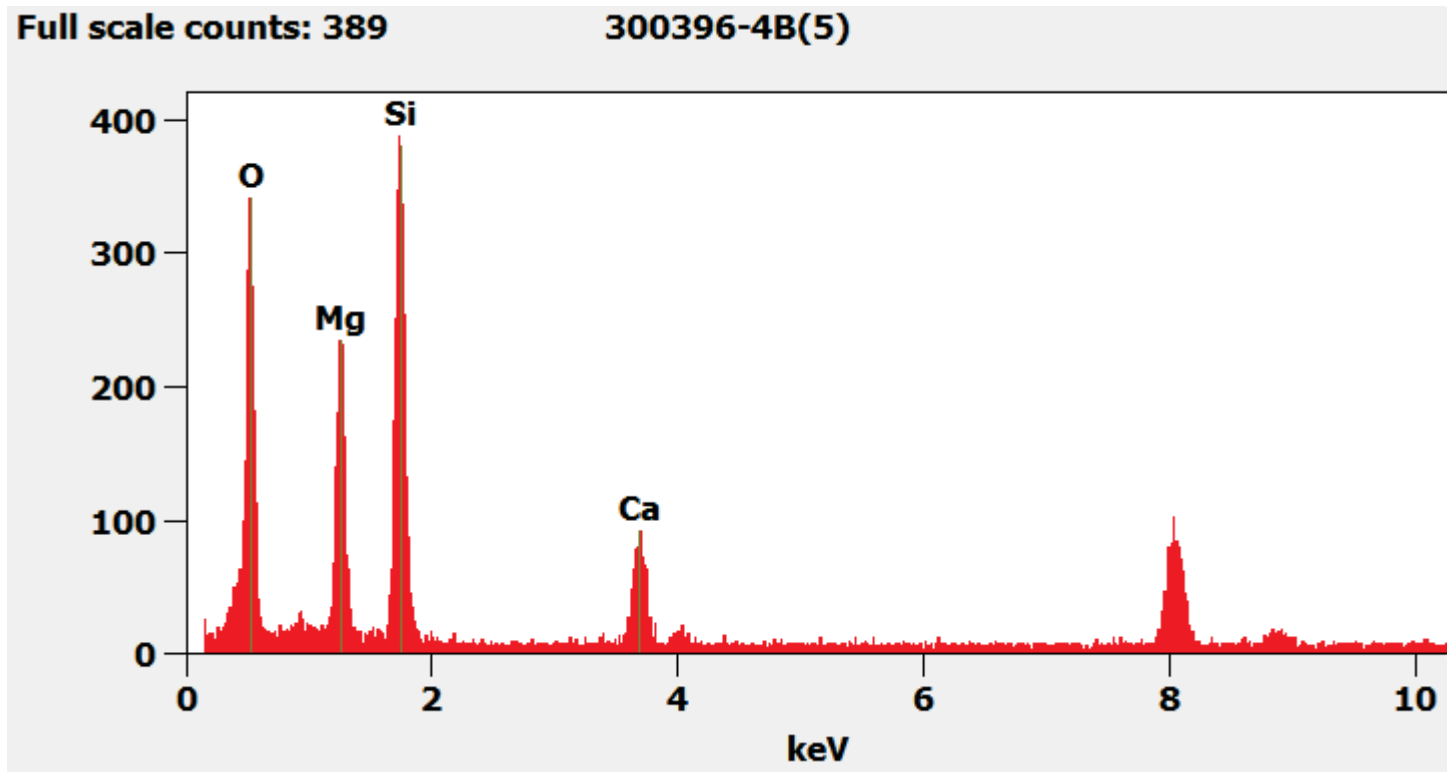
1 µm
HV=100kV
Direct Mag: 1900 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

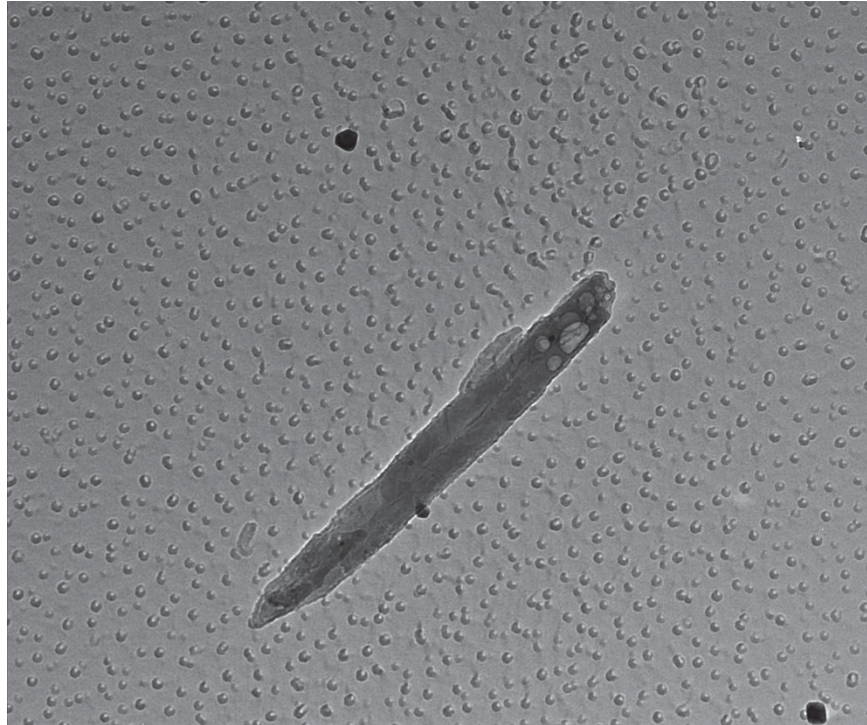
Sample 300396-4B Diffraction pattern from tremolite fiber pictured above.



Sample 300396-4B Chemistry from tremolite fiber pictured above.



Sample 300396-4B Chrysotile structure #1



300396 FDA_108.jpg
Chrysotile Fiber
Cal: 0.541520 nm/pix
10:46 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 nm
HV=100kV
Direct Mag: 19000 x
AMA Analytical Services, Inc

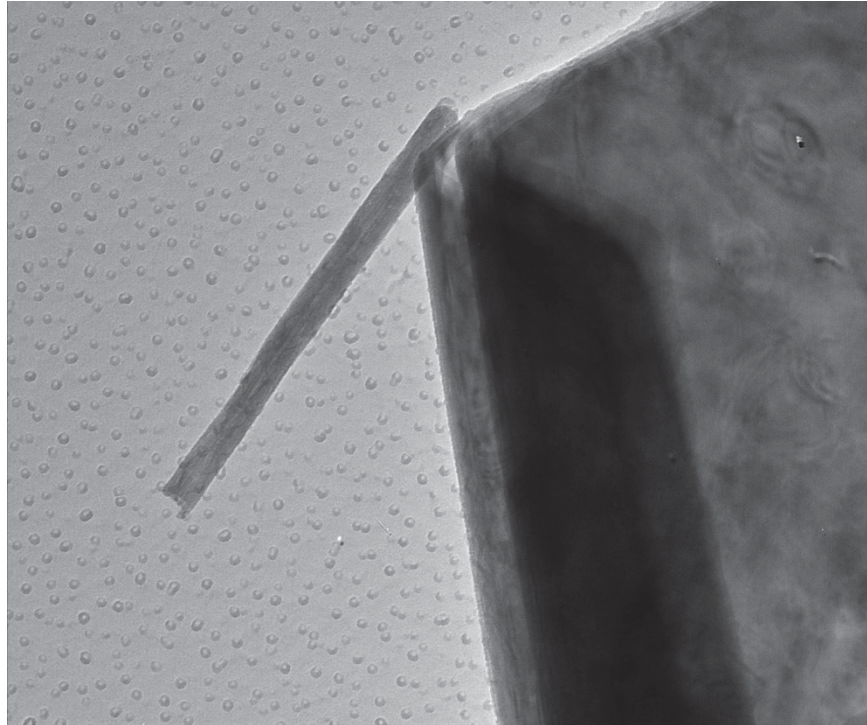
Sample 300396-4B Diffraction pattern from chrysotile structure #1 pictured above



300396 FDA_107.jpg
Chrysotile Diffraction
10:45 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 std. frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Sample 300396-4B Chrysotile structure #2



300396 FDA_110.jpg
Chrysotile Fiber 2
Cal: 0.541520 nm/pix
10:49 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 nm
HV=100kV
Direct Mag: 19000 x
AMA Analytical Services, Inc

Sample 300396-4B Diffraction pattern from chrysotile structure #2 pictured above.

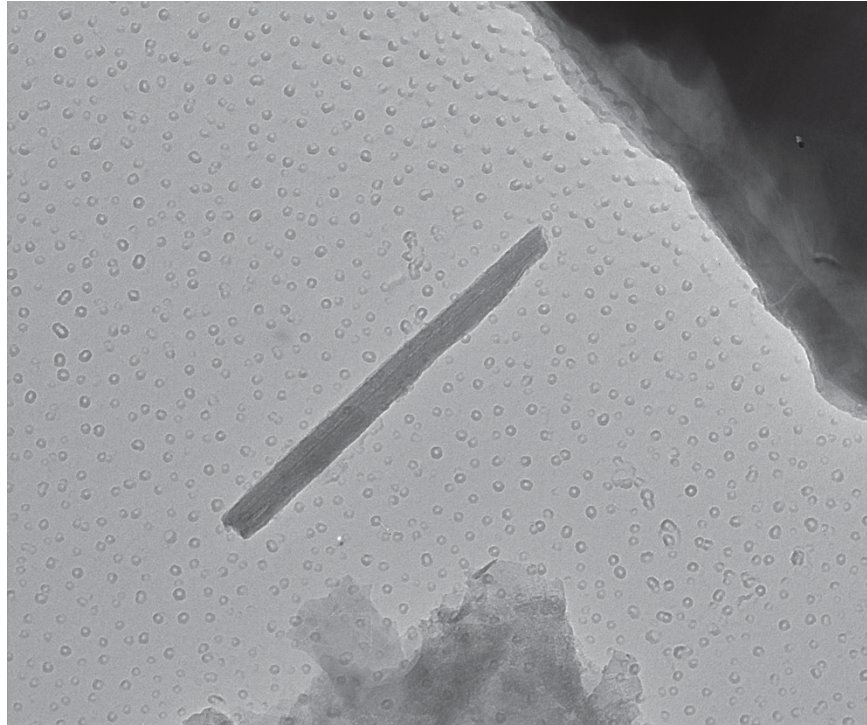


300396 FDA_109.jpg
Chrysotile Diff 2
10:48 4/16/2019
Microscopist: MG

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4B Chrysotile structure #3



300396 FDA_114.jpg
Chry Fiber 3
Cal: 0.541520 nm/pix
11:03 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 nm
HV=100kV
Direct Mag: 19000 x
AMA Analytical Services, Inc

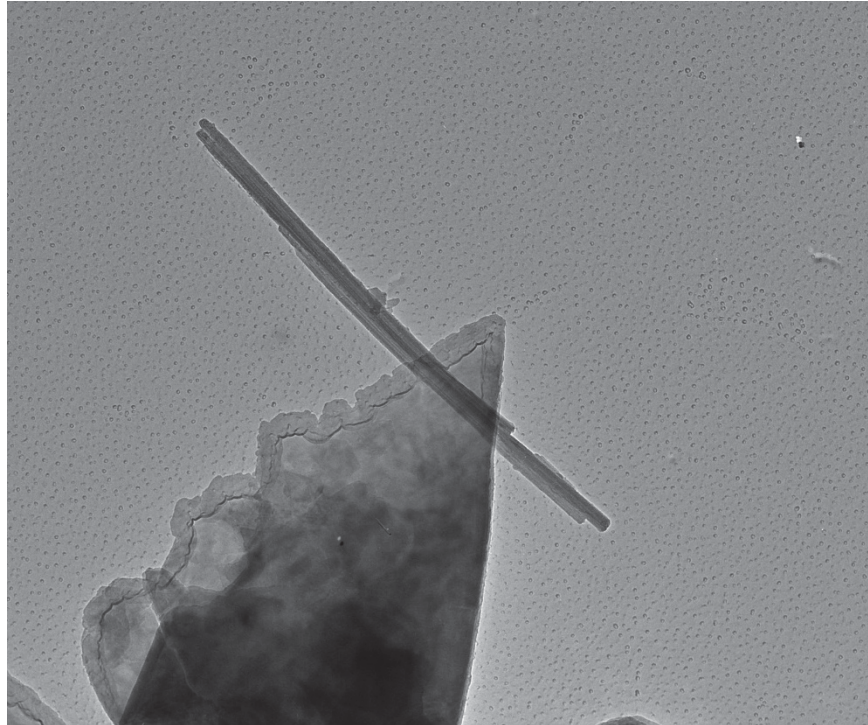
Sample 300396-4B Diffraction pattern from chrysotile fiber #3 pictured above.



300396 FDA_113.jpg
Chry Diff 3
11:02 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Sample 300396-4B Chrysotile structure #5



300396 FDA_119.jpg
Chry Fiber 5
Cal: 0.001429 $\mu\text{m}/\text{pix}$
11:52 4/16/2019
Microscopist: MG
Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

400 nm
HV=100kV
Direct Mag: 7200 x
AMA Analytical Services, Inc

Sample 300396-4B Diffraction pattern for Chrysotile structure #5 pictured above.

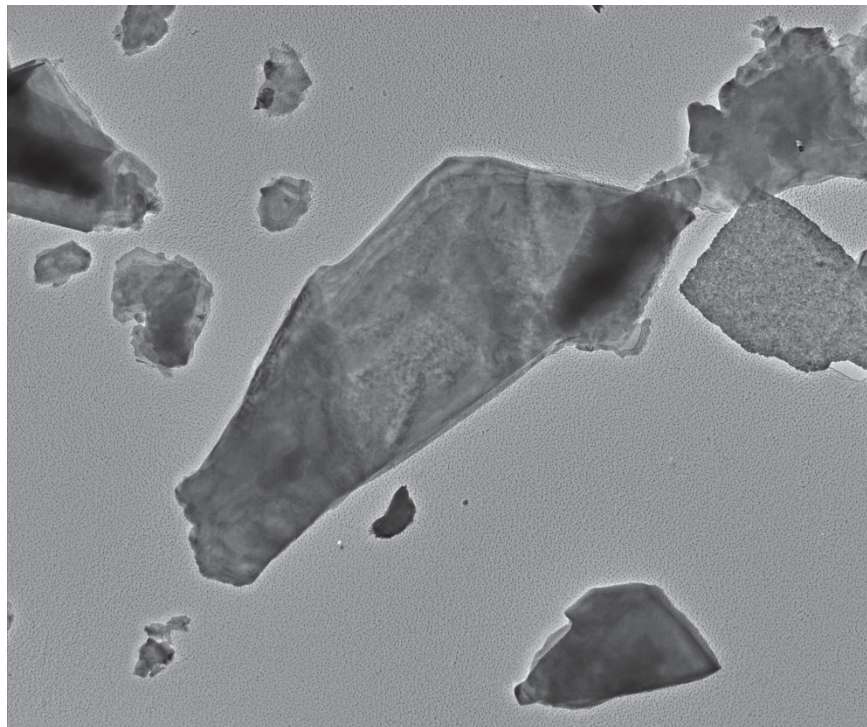


300396 FDA_118.jpg
Chry Diff 5
11:51 4/16/2019
Microscopist: MG

100 (1/Å)
HV=100kV
Cam Len: 0.2200 m
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

Sample 300396-4 Mica particle

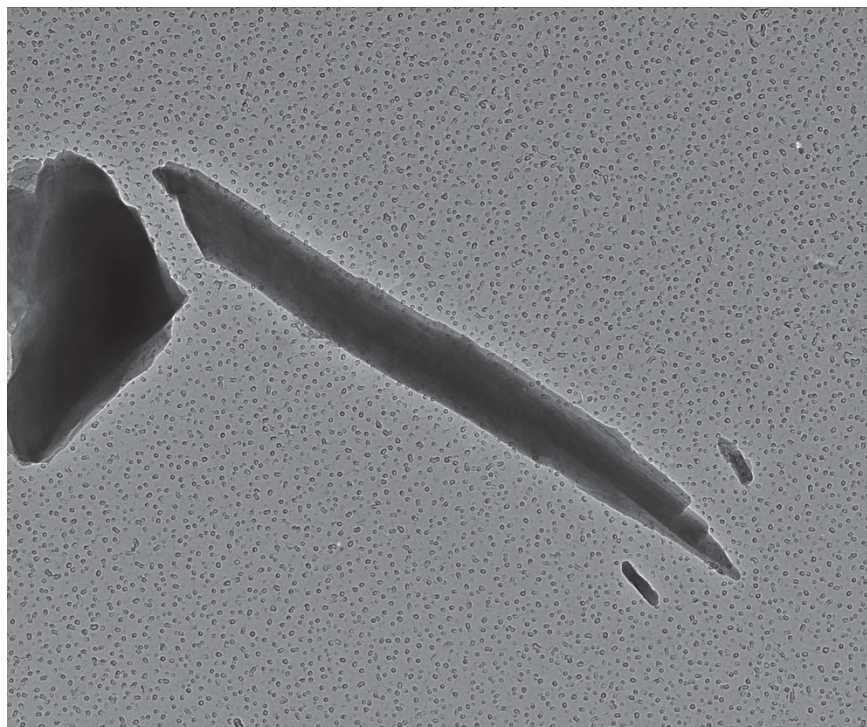


300396 FDA_066.jpg
Mica Particle
Cal: 0.005415 $\mu\text{m}/\text{pix}$
10:30 4/11/2019

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

1 μm
HV=100kV
Direct Mag: 1900 x
AMA Analytical Services, Inc

Sample 300396-4 Talc fiber

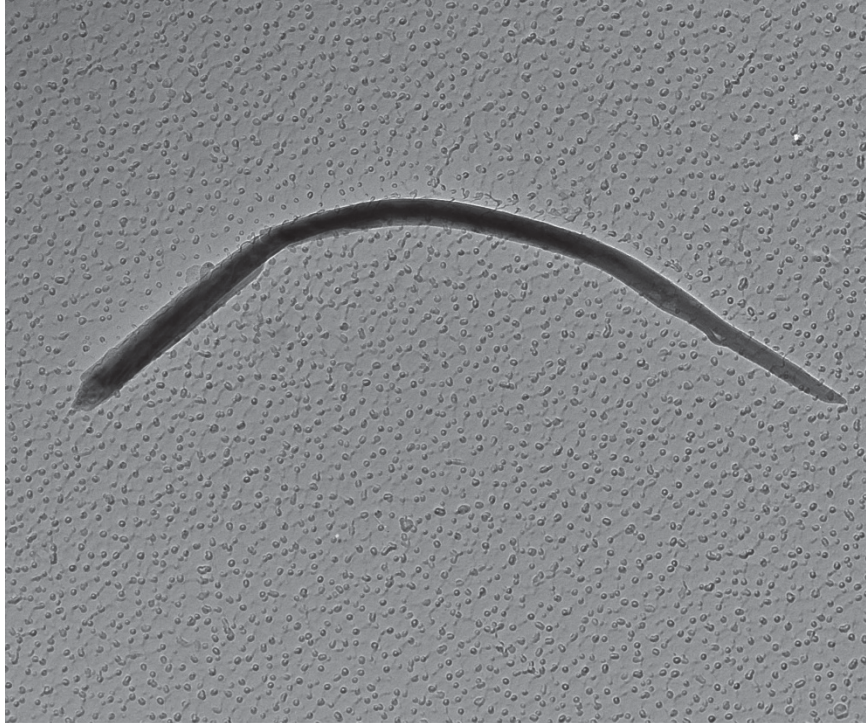


300396 FDA_070.jpg
Talc Fiber
Cal: 0.001429 $\mu\text{m}/\text{pix}$
10:39 4/11/2019

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

400 nm
HV=100kV
Direct Mag: 7200 x
AMA Analytical Services, Inc

Sample 300396-4 Talc ribbon



300396 FDA_076.jpg
Talc Ribbon
Cal: 0.001029 µm/pix
11:10 4/11/2019

200 nm
HV=100kV
Direct Mag: 10000 x
AMA Analytical Services, Inc

Camera: NANOSPRT5, Exposure: 800 (ms) x 5 drift frames, Gain: 1, Bin: 1
Gamma: 1.00, No Sharpening, Normal Contrast

(b) (4)

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