

# Discussion of Potential Changes to ARB Test Method 435:

# Determination of Asbestos Content of Serpentine Aggregate

January 24, 2008, Workshop

Operations Planning & Assessment Section Quality Management Branch Monitoring and Laboratory Division

# Workshop Agenda

- Introduction
- Interlaboratory Study (ILS) Presentation
  - Questions & Answers
- Potential Revisions to Test Method 435 (M435)
  - Processing Procedures
  - Analytical Procedures
  - Laboratory Accreditation
- Revision Schedule/Next Workshop

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## M435 Revision Schedule

- January 24, 2008, Workshop (1st)
  - Rationale & identification of areas of M435 currently being examined for revision
- May/June 2008 Workshop(s)
  - More focused proposed revisions to M435
- August/September Workshop(s)
  - Proposed draft language available for comment
- February 2009 Board Hearing

## **ARB Asbestos Regulations**

- 1986 Identified asbestos as an airborne toxic contaminant (TAC)
- 1991 Airborne Toxic Control Measure (ATCM) for Surfacing Applications
   Revised 2001
- 1991 M435: Determination of Asbestos Content of Serpentine Aggregate
- 2002 ATCM for Construction, Grading, Quarrying, and Surface Mining Operations
- ➤ M435 referenced in both ATCMs

#### Rationale for M435 Revision

- Observations by parties regarding the variability of laboratory equipment, M435 procedures, and analytical results
- Results of ARB ILS show that certain M435 laboratory practices result in differences in the % asbestos reported.

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## Preliminary Interpretation of the Interlaboratory Study for ARB Test Method 435:

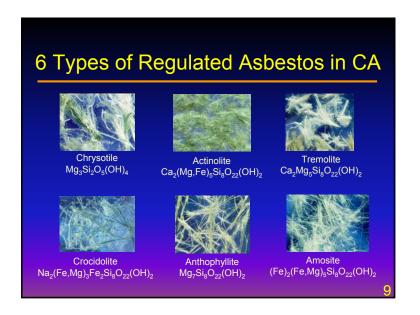
Determination of Asbestos Content of Serpentine Aggregate

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## **ARB Test Method 435**

- Airborne Toxic Control Measure (ATCM) for Surfacing Applications
  - <0.25% asbestos content limit of aggregate material
- ATCM for Construction, Grading, Quarrying, and Surface Mining Operations
  - Dust control measures required in areas with ≥0.25% asbestos
- Used by Dept. of Toxic Substances Control (DTSC) to determine asbestos content of soils at new school construction sites in CA;
- Used nationwide as a bulk method to determine asbestos content in soils.

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Observations by some parties regarding Test Method 435 (M435):

- "Guidelines in Test Method 435 allow laboratories great latitude in processing and analytical procedures;"
- "Laboratories prepare and analyze soil and rock samples in different ways;"
- "Different laboratories obtain differing results when analyzing the same samples for asbestos."

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# Interlaboratory Study (ILS) Objectives

- To investigate variability in preparation and analytical procedures used by laboratories applying M435;
- To determine whether these differences affect asbestos content determination.

Test Method 435 Protocol

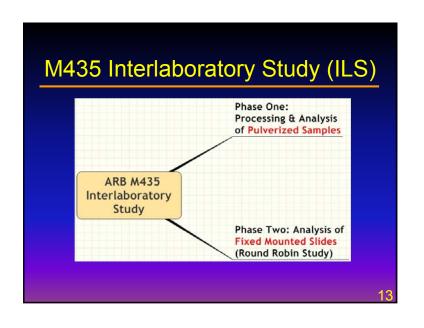
Geologic Sample

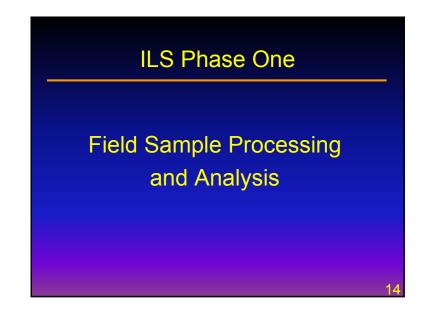
Dry, crush to <3/8" nominal diameter, reduce to 1 pint aliquot

Pulverize majority to <75 µm diameter

Analyze using Polarized Light Microscopy (PLM)

400-point count rules Determine % Asbestos





## **Phase One Objectives**

- To obtain qualitative information on variability of equipment, sample processing, protocols, and fibercounting practices among laboratories;
- To determine whether these differences result in variations of reported % asbestos.

Phase One:
Crushing of Field Sample

ARB archived 1 gal.
NOA sample

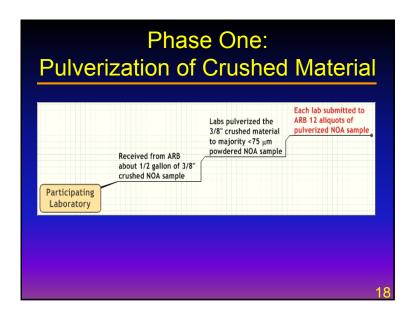
ARB supervised 3/8" crushing asbestos (NOA) sample
ARB supervised 3/8" crushing a homogenization of 4 gals.
NOA sample

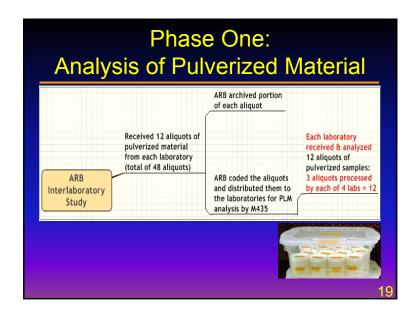
ARB darchived 2 gals.
of 3/8" crushed sample a homogenization of 4 gals.
NOA sample

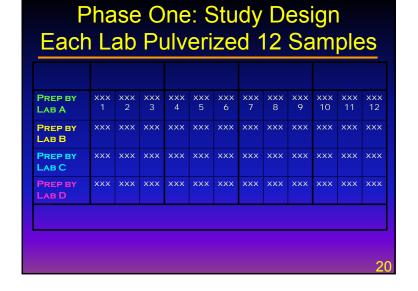
ARB supervised 3/8" crushing a sample a homogenization of 4 gals.
NOA sample

ARB darchived 2 gals.
of 3/8" crushing a homogenization of 4 gals.
ARB distributed 1/2 gallon crushed sample to each laboratory for pulverization









# Phase One: Study Design Each Lab Analyzed 12 Samples

	ANA Lab	LYSE:	S ВҮ	Ana Lab	LYSE! B	S BY	Ana Lab	LYSE:	S BY	Ana Lab	LYSE! D	S ВҮ
PREP BY LAB A	xxx 1	xxx 2	xxx 3	XXX	XXX	XXX	XXX	XXX	xxx	XXX	XXX	XXX
PREP BY LAB B	xxx 4	xxx 5	xxx 6	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx
PREP BY LAB C	xxx 7	xxx 8	xxx 9	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx
PREP BY LAB D	xxx 10	xxx 11	xxx 12	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx	xxx

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## Phase One: Coding of Results

To avoid attribution of the study results to any specific Participating Laboratory, the following tables and graphs use letter names to refer to the laboratories.

These letter names are for discussion references only, and have no continuity in the presentation.

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# Phase One: Analytical Results

PREP BY	0.75										
Lab E	0.75			0.0*	0.0*	0.0*	0.75	1.00	0.75		
PREP BY LAB F	1.00		1.00	0.0*	0.0*	0.0*	0.25	0.50	0.50		
PREP BY LAB G	0.0*		0.25	0.0*	0.0*	0.0*	0.0*		0.0*		
PREP BY LAB H	0.75	1.00		0.0*	0.25	0.0*		0.50	1.00		

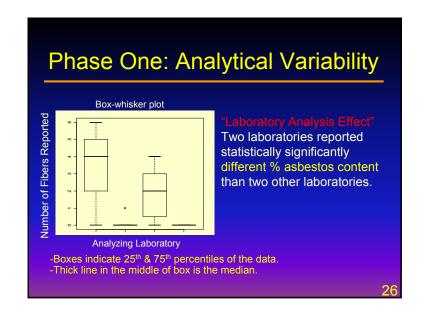
## Phase One: Analytical Results

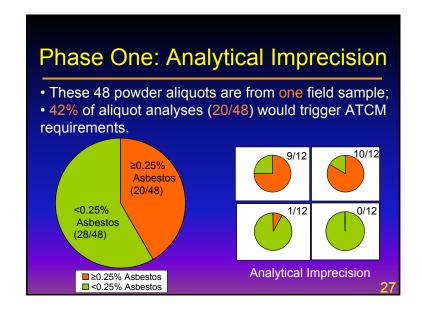
	Ana Lab	LYSES A	S BY	Ana Lab	LYSES B	BY	Ana Lab	LYSES C	BY	Ana Lab	LYSES D	BY
PREP BY LAB E	0.75	1.25	1.25	0.0*	0.0*	0.0*	0.75	1.00	0.75	0.00	0.00	0.00
PREP BY LAB F	1.00	1.50	1.00	0.0*	0.0*	0.0*	0.25	0.50	0.50	0.00	0.00	0.00
PREP BY LAB G	0.0*	0.0*	0.25	0.0*	0.0*	0.0*	0.0*	0.0*	0.0*	0.00	0.00	0.00
PREP BY LAB H	0.75	1.00	1.50	0.0*	0.25	0.0*	0.75	0.50	1.00	0.00	0.00	0.00
Mhen "<0.25%" o	lote: Whe										ed for thi	s table.

- Labs A and C reported asbestos in the majority of aliquots;
- · Labs B and D reported very little asbestos.

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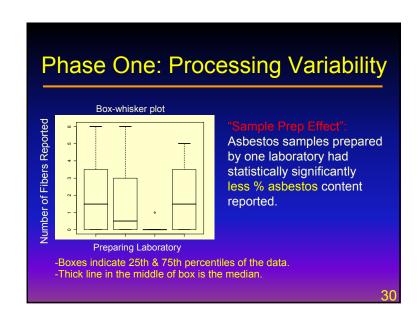
	Ana Lab	LYSES A	S ВҮ	Ana Lab	LYSE! B	S ВҮ	Ana Lab	LYSE!	БВҮ	LYSES D	
PREP BY Lab E	0.75	1.25	1.25	0.0*	0.0*	0.0*	0.75	1.00	0.75		
PREP BY LAB F	1.00	1.50	1.00	0.0*	0.0*	0.0*	0.25	0.50	0.50		
PREP BY LAB G	0.0*	0.0*	0.25	0.0*	0.0*	0.0*	0.0*	0.0*	0.0*		
PREP BY LAB H	0.75	1.00	1.50	0.0*	0.25	0.0*	0.75	0.50	1.00		





	Ana Lab	LYSES A	BY	Ana Lab	LYSES B	BY	Ana Lab	LYSES C	BY	Ana Lab	LYSES D	ву
PREP BY Lab E	0.75	1.25	1.25	0.00	0.00	0.00	0.75	1.00	0.75	0.00	0.00	0.00
PREP BY LAB F	1.00	1.50	1.00	0.00	0.00	0.00	0.25	0.50	0.50	0.00	0.00	0.00
PREP BY LAB G	0.00	0.00	0.25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
PREP BY LAB H	0.75											

	Ana Lab	LYSE! A	S BY	Ana Lab	LYSE! B	s вү	Ana Lab	LYSES C	S ВҮ	Ana Lab	LYSE! D	S ВҮ
PREP BY Lab E	0.75	1.25	1.25	0.0*	0.0*	0.0*	0.75	1.00	0.75	0.00	0.00	0.00
PREP BY Lab F	1.00	1.50	1.00	0.0*	0.0*	0.0*	0.25	0.50	0.50	0.00	0.00	0.00
PREP BY Lab G	0.0*		0.25									
PREP BY LAB H	0.75	1.00	1.50	0.0*	0.25	0.0*	0.75	0.50	1.00	0.00	0.00	0.00



# Phase One: Processing Variability

- Were there any noticeable differences among the pulverized aliquots in Phase One?
- If so, what were these differences?
- Did these differences affect the % asbestos reported?

Prepared by Lab I

Prepared by Lab I

Prepared by Lab J

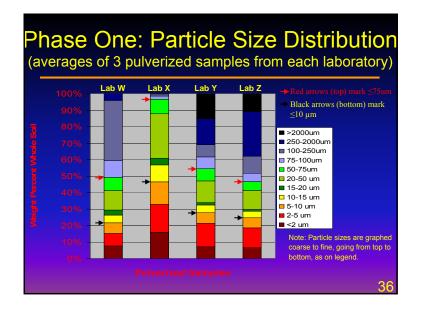
# Phase One: Processing Variability Quantitative Analysis

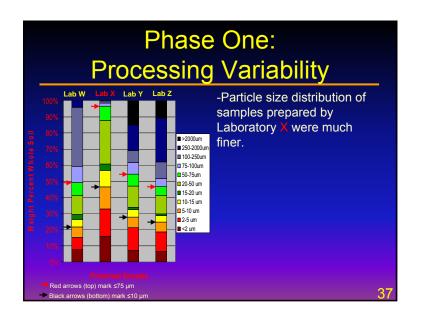
 Did laboratories produce pulverized aliquots with similar particle size distribution?

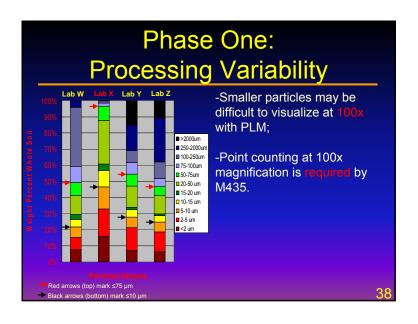
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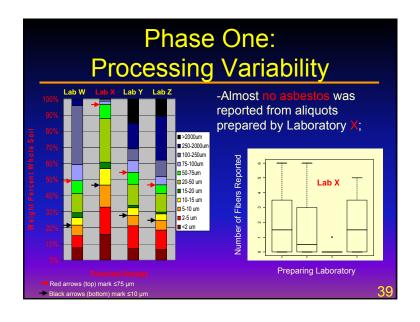
# Phase One: Particle Size Distribution (3 aliquots from each laboratory) Particle size analysis (PSA) by pipette sand 50-2000μm silt 2-50μm <5μm, <10μm, <15μm, <20μm clay <2μm Dry sieving 50μm, 75μm (200 mesh), 100μm, 250μm, 500μm, 1000μm, 2000μm, >2000μm

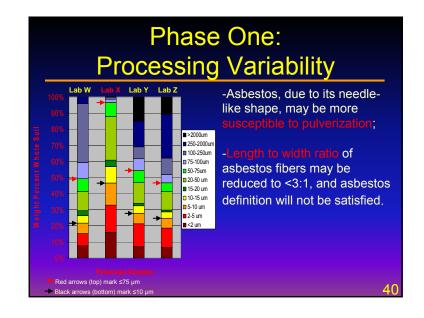
#### Phase One: Lab M **PSA Variability** nnl Three aliquots from Lab N each laboratory; Particle size analyses (PSA) Lab O show differences among laboratories 60 Lab P in % mass of size fractions: · Labs O and P have 22 um 2-50 um 50-2000 um 32000 um particles >2000um. Particle Size











# Phase One: Processing Variability Quantitative Analysis

 Did laboratories produce pulverized aliquots with similar particle size distribution?

No, different methods of pulverization produced different particle size distributions.

 Very fine particle size distribution appears to significantly decrease the % asbestos reported.

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## **ILS Phase Two**

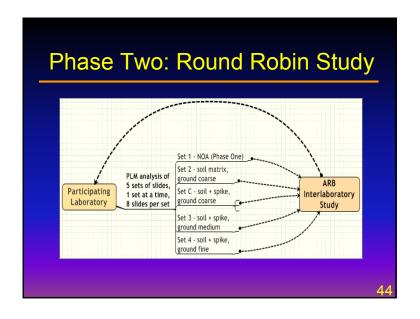
Analysis of Fixed Mounted Slides

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# Phase Two Objectives

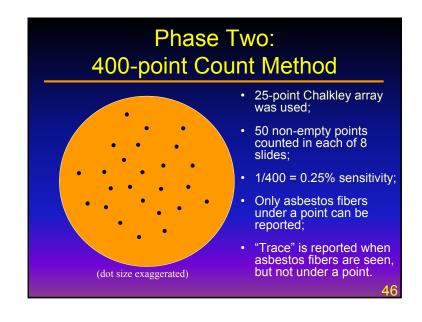
- To observe variability between laboratories in asbestos sample analysis, while minimizing sample processing effects;
- •To observe the effect of counting rules on number of asbestos fibers reported;
- •To observe the effect of sample particle size distribution on number of asbestos fibers reported;
- •To observe variability among laboratories quantifying naturally-occurring asbestos (NOA) versus NIST\* standard reference asbestos.

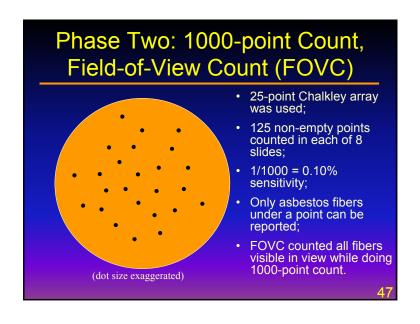
\*NIST - National Institute of Standards and Technology

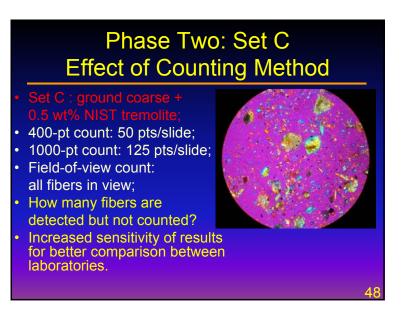


#### Phase Two: 5 Sets of Fixed Slides + Ground Samples Description **Objectives** Set 1 NOA sample, NOA vs. NIST\* previously analyzed in Phase 1 reference asbestos Set 2 Soil matrix, no spike, Assess asbestos content of soil matrix ground co Soil matrix + 0.5 wt% NIST\* tremolite, Compare asbestos content using 400-point ct, 1000-point ct, field-of-view ct 3 counting methods Set 3 Soil matrix + 0.5 wt% NIST tremolite, Assess effect of ground medium particle size Set 4 Soil matrix + 0.5 wt% NIST tremolite, distribution on ground fine asbestos count

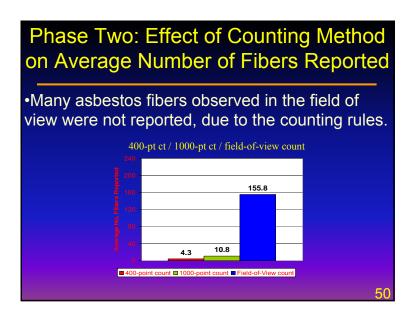
\* NIST - National Institute of Standards and Technology

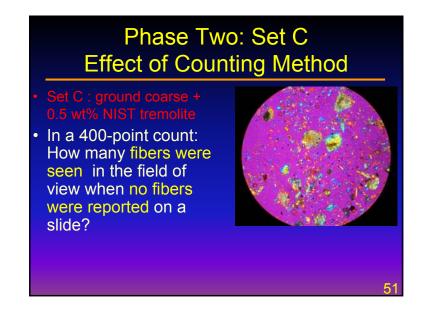


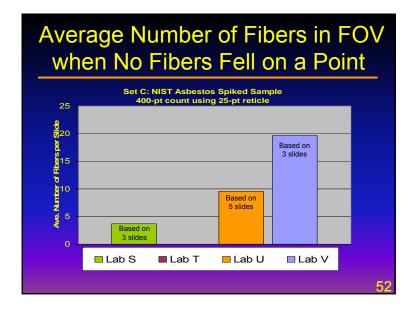




		Lab W	Lab X	Lab Y	Lab Z	Ave. No. of Fibers
et C 00-point ount	# of fibers	6	0	4	7	
Set C 000-pt count	# of fibers	8	0	9	26	
Set C Field-of- /iew count	# of fibers	53	0	205	365	







## Phase Two: Effect of Particle Size

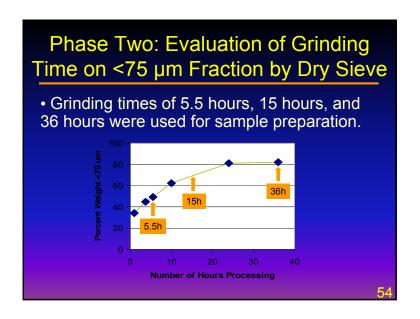
 Compared the number of asbestos fibers reported in a 400-point count analysis of:
 Set C – ground coarse with 0.5 wt% NIST

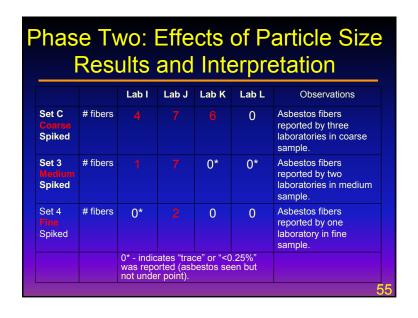
Set C – ground coarse with 0.5 wt% NIST tremolite;

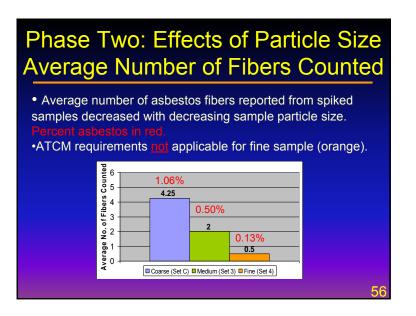
Set 3 – ground medium with 0.5 wt% NIST tremolite;

Set 4 – ground fine with 0.5 wt% NIST tremolite.

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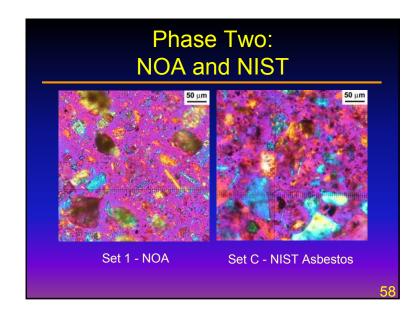






# Phase Two: NOA and NIST Asbestos

- Set 1 NOA sample from Phase One;
- Set C, Set 3, Set 4 spiked with NIST tremolite asbestos;
- 400-point count analysis.



Phase To Number of F				
	Lab M	Lab N	Lab O	Lab P
Set 1 NOA	0	0*	35	0*
Set 2 Soil Matrix	0	0	5	0
Set C – 0.5 wt% spike NIST tremolite	0	4	7	6
•NOA reported by one I -Another lab reporte set of 12 aliquots; •NIST asbestos reporte	aboratory d NOA in	Phase Or	ne from the	t under a point
				59

	se One				
		Lab Q	Lab R	Lab S	Lab T
Phase One 400-pt count NOA	Sum of all fibers, 12 aliquots	1	41	24	0
Phase Two 400-pt count NIST tremolite + NOA	Sum of all fibers, Sets C, 3, 4 + Set 1, 2	6	61	5	0
	Totals	7	102	29	0
					(

## Some Conclusions from ILS

- Laboratories use different processing equipment and protocols;
  - Result in varying particle size distribution of samples;
- Finer particle size distribution is one factor resulting in lower % asbestos reported;
  - Observed in Phase One:
    - Preparation Effect
  - Observed in Phase Two:
    - Set C (coarse) vs. Set 4 (fine)

Some Conclusions from ILS

 Sampling using 400-point count analysis lowers reportable number of asbestos fibers by 1 to 2 orders of magnitude when compared to the field-of-view count;

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## Some Conclusions from ILS

- Fiber identification criteria are not uniform among laboratories;
  - Phase One: Laboratory Analysis Effect
  - Phase Two: Laboratories reported a wide range of number of asbestos fibers
  - One laboratory did not detect NOA nor report NIST standard reference asbestos.

#### **ARB Future Activities**

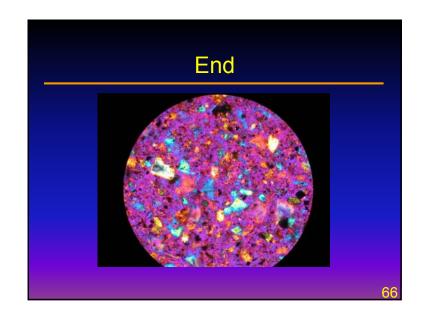
- To work with stakeholders in identifying variables that can reduce laboratory processing and analytical variability when applying Test Method 435.
- To revise Test Method 435 accordingly.

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## Acknowledgment

- Asbestos TEM Laboratories, Inc.
- EMSL Analytical, Inc.
- Forensic Analytical Laboratories, Inc.
- R J Lee Group, Inc.
- NOA Laboratory Working Group, SAGE of El Dorado County
- Southard Laboratory, Dept. of LAWR, University of California - Davis

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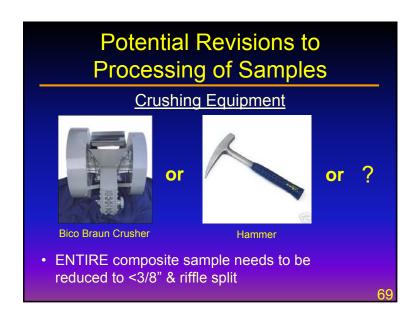
## Workshop Agenda

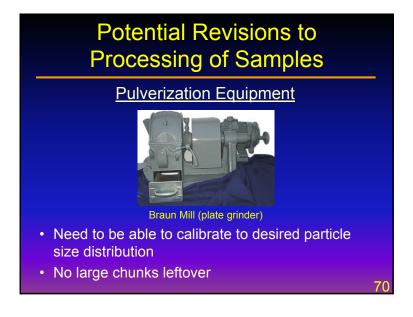
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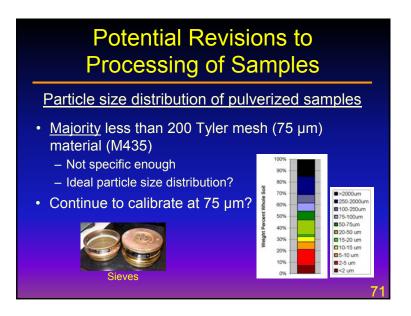
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# Potential Revisions to Analytical Procedures

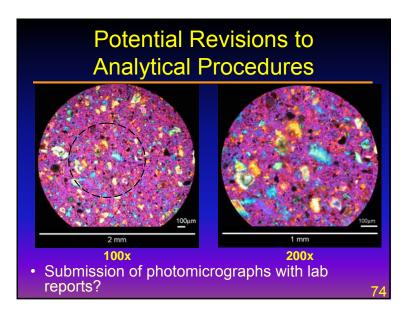
#### Magnification

- Specify magnification during identification
  - M435 currently silent
- Increase magnification during point count
  - morphology more easily determined
  - reticle dot becomes "smaller"

#### **Increase in Point Count**

- Would lower detection limit below ATCM's "trigger" level
- Would keep spatial representativeness intact with increase in magnification

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# Potential Revisions to Analytical Procedures

- Reticles
  - Unbiased sampling?
- Mechanical Stage
  - Predetermined movement of crosshair reticle



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## Potential Revisions to Analytical Procedures

#### Fiber Identification / Definition

- Labs have different interpretation of what should be identified as asbestos
- M435 criteria need to be more explicit
- ARB will work with OEHHA to ensure revised definition reflects health information

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# **Laboratory Accreditation Promotes**

- Measurement accuracy
- Accepted quality control and good laboratory practices
- Quality assessment through proficiency testing
- Corrective action for nonconformities
- ➤ No current accreditation requirement to perform M435 analysis

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### Need for M435 Accreditation

- ILS showed significant variability in M435 results among labs accredited for PLM analysis of bulk asbestos products;
- Current accreditation programs are not specific to naturally-occurring asbestos (NOA) samples
  - NVLAP/AIHA accreditation for PLM use an EPA method for the analysis of asbestos in building materials (EPA600/R-93/116);

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# Laboratory Accreditation Organizations

- NVLAP National Voluntary Laboratory Accreditation Program
- NELAP/ELAP National Environmental Laboratory Accreditation Program / California ELAP within the California Department of Public Health
- AIHA American Industrial Hygiene Association

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## M435 Revision Schedule

- January 24, 2008, Workshop (1st)
  - Rationale & identification of areas of M435 currently being examined for revision
- May/June 2008 Workshop(s)
  - More focused proposed revisions to M435
- August/September Workshop(s)
  - Proposed draft language available for comment
- February 2009 Board Hearing

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## Workshop Two

- Time frame: May/June 2008;
- Possible venues?

