

The background of the slide features a stylized city skyline at the bottom, composed of various geometric shapes in shades of blue and white. Above the skyline, the background is a solid blue color with several stylized, light blue clouds scattered across it. The title text is centered in the upper half of the slide.

Detection Limits for Source Emissions Sampling Methods: The Uncertainty of Uncertainty.

Sheri Heldstab
CHESTER LabNet

Water Sample

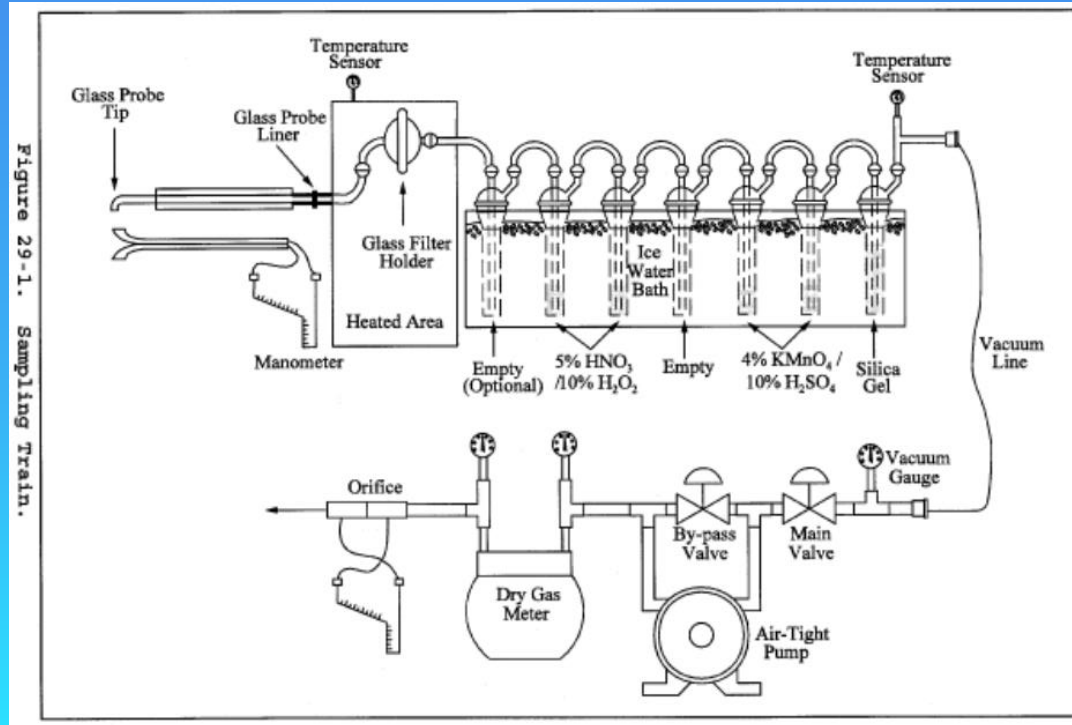


Air Sample

Collecting Water Samples

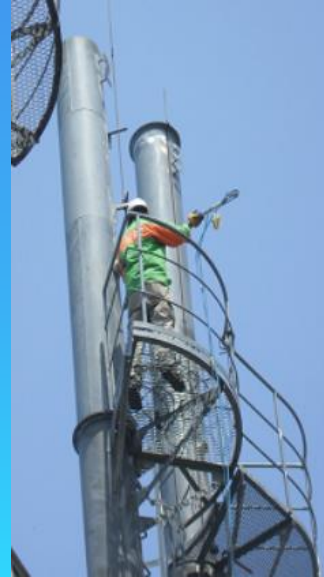


Collecting Source Emission Samples[‡]



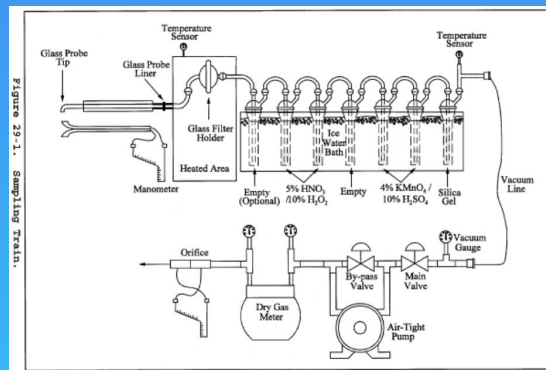
[‡] 40 CFR Part 60, Appendix A-8, Method 29

Collecting Source Emission Samples





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Collect Sample

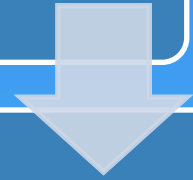


Laboratory Analysis



Data Reduction and Modeling

Collect Sample



Laboratory Analysis



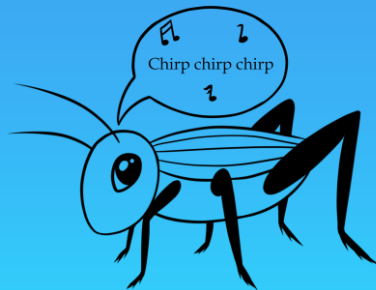
Data Reduction and Modeling

Source Sampling Collection Uncertainties

- Glassware preparation
- Analyzer drift
- Accuracy of O₂/CO₂ measurements (in turn affects your calculation of molecular weight, sample volume, flow, etc.)
- Experience/skill of testers/Human error
- Quality of reagents
- Environmental conditions
- Source stream homogeneity
- Sample loss due to leaks
- Measurements of pressure and temperature
- Thermocouples
- Number of points/port used
- Size/alignment of the nozzle during sampling (straight into the flow?)
- Flow meter uncertainty
- Leak during run (2+ hours continuous)
- Sample bottle type and cleanliness
- Interfering gases
- Field balances and other standards (field balance, field caliper, field barometer, etc.)
- Flow measurements (many factors go into this alone)
- Quality of gas standards
- Measurements of pressure and temperature
- Length of sample run(s) (what snapshot of the process are you capturing?)
- Number of runs (gives you some sense of repeatability)
- Pitot specifications
- Sampling location
- Moisture content of gasses (impinger pH)
- Meter volume
- Recovery of sample in the field (cleanliness?)
- Post-analysis calculations to lbs/year or...

Source Sample Collection Uncertainties

Where are these accounted for in the Detection Limit?



Collect Sample



```
graph TD; A[Collect Sample] --> B[Laboratory Analysis]; B --> C[Data Reduction and Modeling];
```

Laboratory Analysis

Data Reduction and Modeling

Laboratory Detection Limits

NELAC accredited labs follow “Definition and Procedure for the Determination of the Method Detection Limit, Revision 2” (aka “MUR method”)

- Standard deviation of low concentration standards.
- Standard deviation of, and concentration of, blank samples.
- Taken through entire process, including all preparatory steps.

Does not take into account sampling activities.

Collect Sample



```
graph TD; A[Collect Sample] --> B[Laboratory Analysis]; B --> C[Data Reduction and Modeling];
```

Laboratory Analysis

Data Reduction and Modeling

Modeling Math!!

Uncertainty
from:

Sampling? No.

Lab? Yes.

Uncertainty
from other
measurements?

No.

At each Exposure Location...

$TAC_1 \text{ emission rate} \times TEU_A \text{ dispersion factor}$

+

$TAC_1 \text{ RBC at Chronic Exposure Location}$

$TAC_2 \text{ emission rate} \times TEU_A \text{ dispersion factor}$

+

$TAC_2 \text{ RBC at Chronic Exposure Location}$

$TAC_2 \text{ emission rate} \times TEU_R \text{ dispersion factor}$

+

$TAC_2 \text{ RBC at Chronic Exposure Location}$

$TAC_3 \text{ emission rate} \times TEU_R \text{ dispersion factor}$

+

$TAC_3 \text{ RBC at Chronic Exposure Location}$

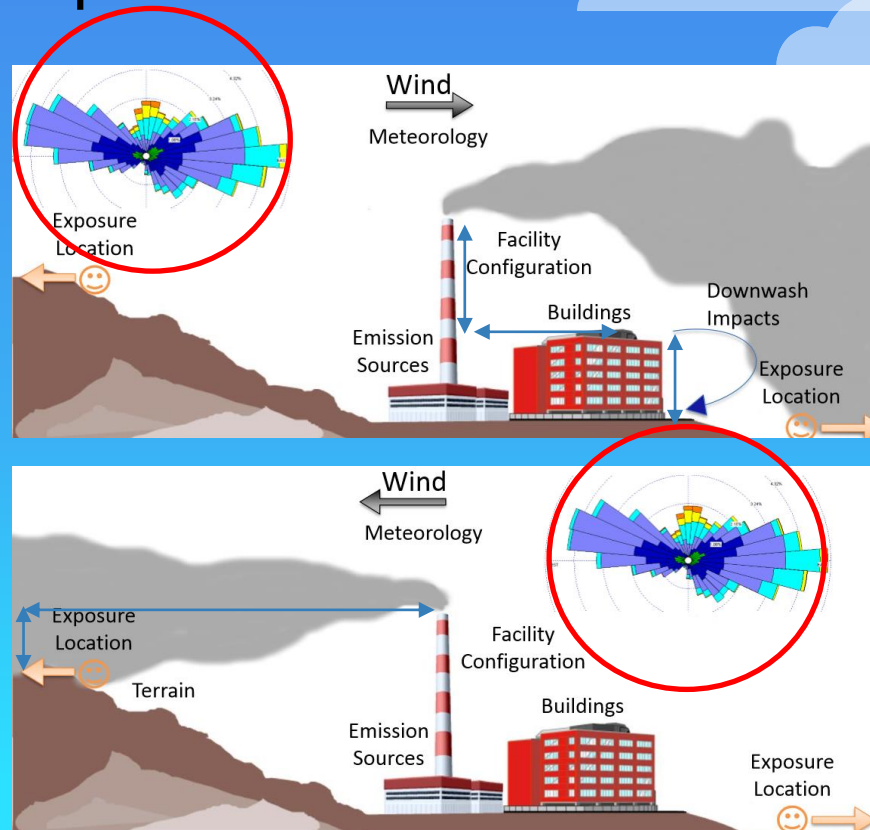
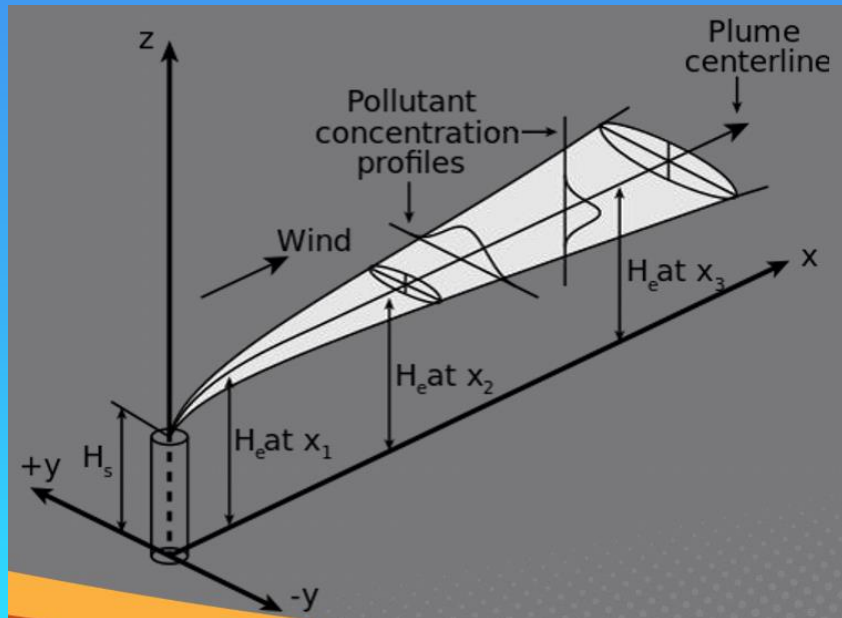
$TAC_4 \text{ emission rate} \times TEU_R \text{ dispersion factor}$

+

$TAC_4 \text{ RBC at Chronic Exposure Location}$

Modeling Step 1: Math!!

Don't worry, this won't be on the test.



Remember This Math?

At each Exposure Location...

$$\begin{aligned}
 & \frac{TAC_1 \text{ emission rate} \times TEU_A \text{ dispersion factor}}{TAC_1 \text{ RBC at Chronic Exposure Location}} + \\
 & \frac{TAC_2 \text{ emission rate} \times TEU_A \text{ dispersion factor}}{TAC_2 \text{ RBC at Chronic Exposure Location}} + \\
 & \frac{TAC_2 \text{ emission rate} \times TEU_R \text{ dispersion factor}}{TAC_2 \text{ RBC at Chronic Exposure Location}} + \\
 & \frac{TAC_3 \text{ emission rate} \times TEU_B \text{ dispersion factor}}{TAC_3 \text{ RBC at Chronic Exposure Location}} + \\
 & \frac{TAC_4 \text{ emission rate} \times TEU_R \text{ dispersion factor}}{TAC_4 \text{ RBC at Chronic Exposure Location}} +
 \end{aligned}$$

Modeling Step 2: Determination of Exposure Locations



- Distances measured by Google Earth. Uncertainty?
- Even if using a laser distance meter, uncertainty is still there (and many other problems).
- May be measured using USGS or other gov't maps, still have uncertainty.

Distance measurements contribute to modeling uncertainty, because...

Remember This Math?

At each Exposure Location...

$$\frac{TAC_1 \text{ emission rate} \times TEU_A \text{ dispersion factor}}{TAC_1 \text{ RBC at Chronic Exposure Location}} +$$

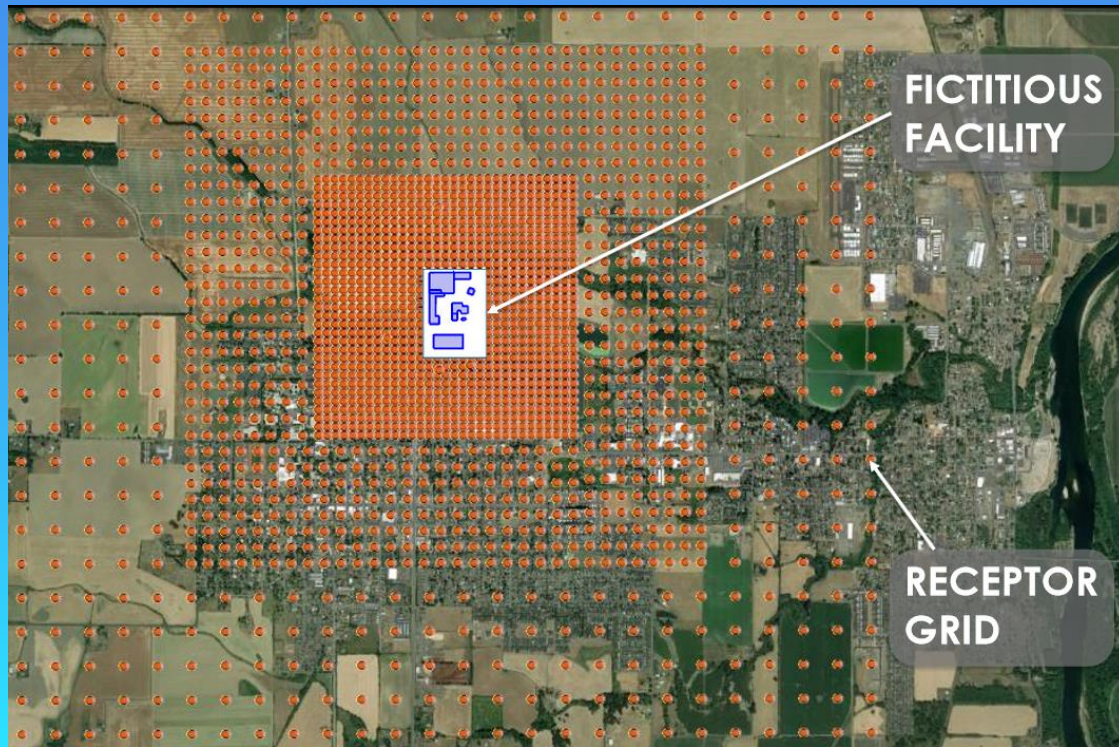
$$\frac{TAC_2 \text{ emission rate} \times TEU_A \text{ dispersion factor}}{TAC_2 \text{ RBC at Chronic Exposure Location}} +$$

$$\frac{TAC_2 \text{ emission rate} \times TEU_B \text{ dispersion factor}}{TAC_2 \text{ RBC at Chronic Exposure Location}} +$$

$$\frac{TAC_3 \text{ emission rate} \times TEU_B \text{ dispersion factor}}{TAC_3 \text{ RBC at Chronic Exposure Location}} +$$

$$\frac{TAC_4 \text{ emission rate} \times TEU_B \text{ dispersion factor}}{TAC_4 \text{ RBC at Chronic Exposure Location}} +$$

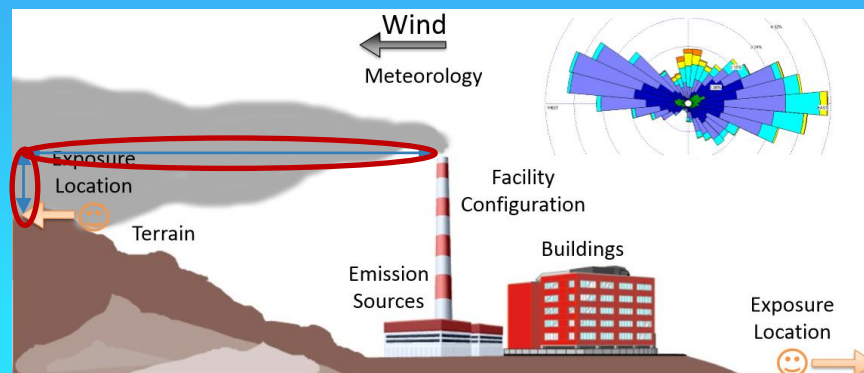
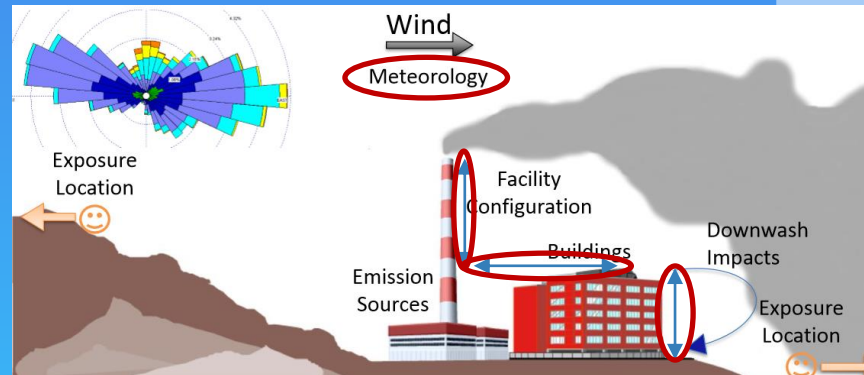
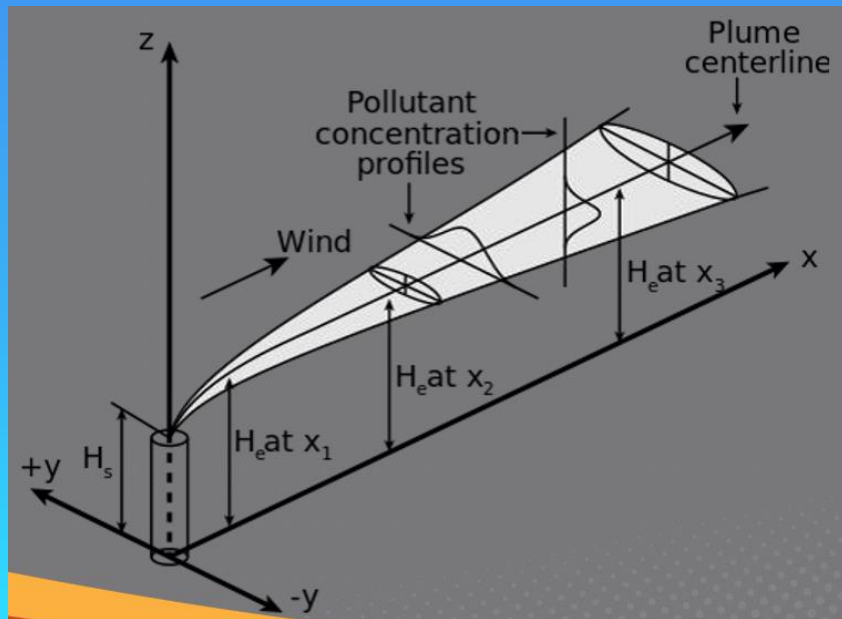
Modeling Step 3: More Math!!



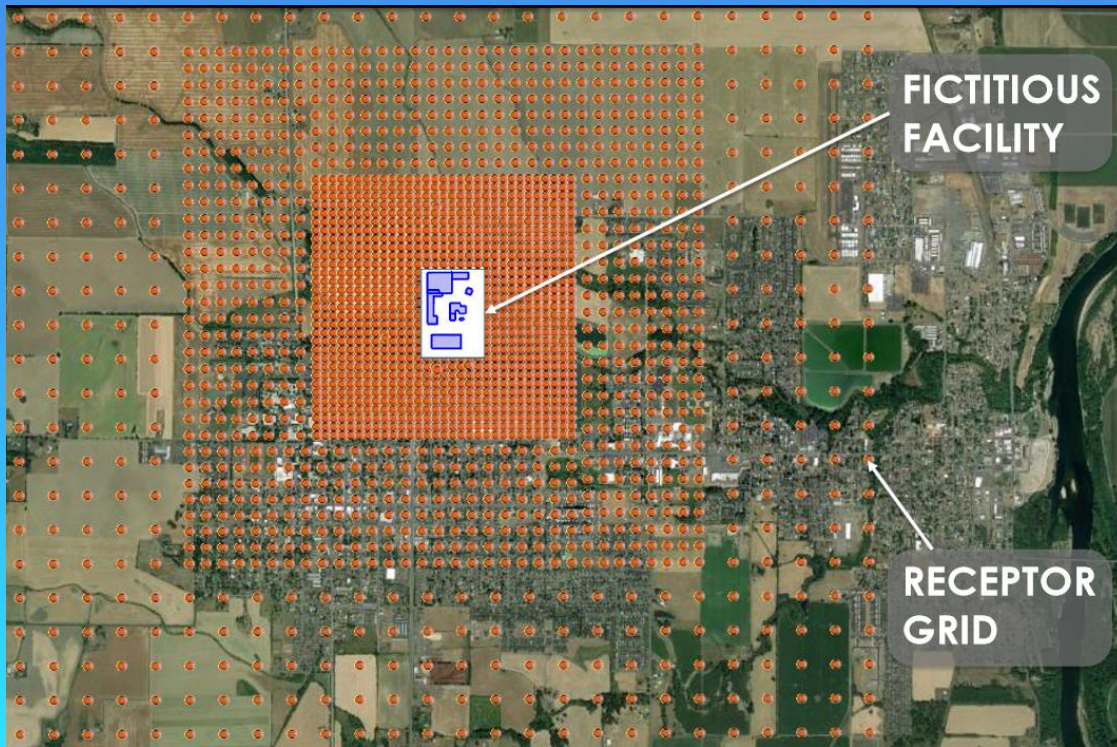
2,149 receptor points
requiring data
reduction using
plume concentrations.

Remember This??

Plume concentrations



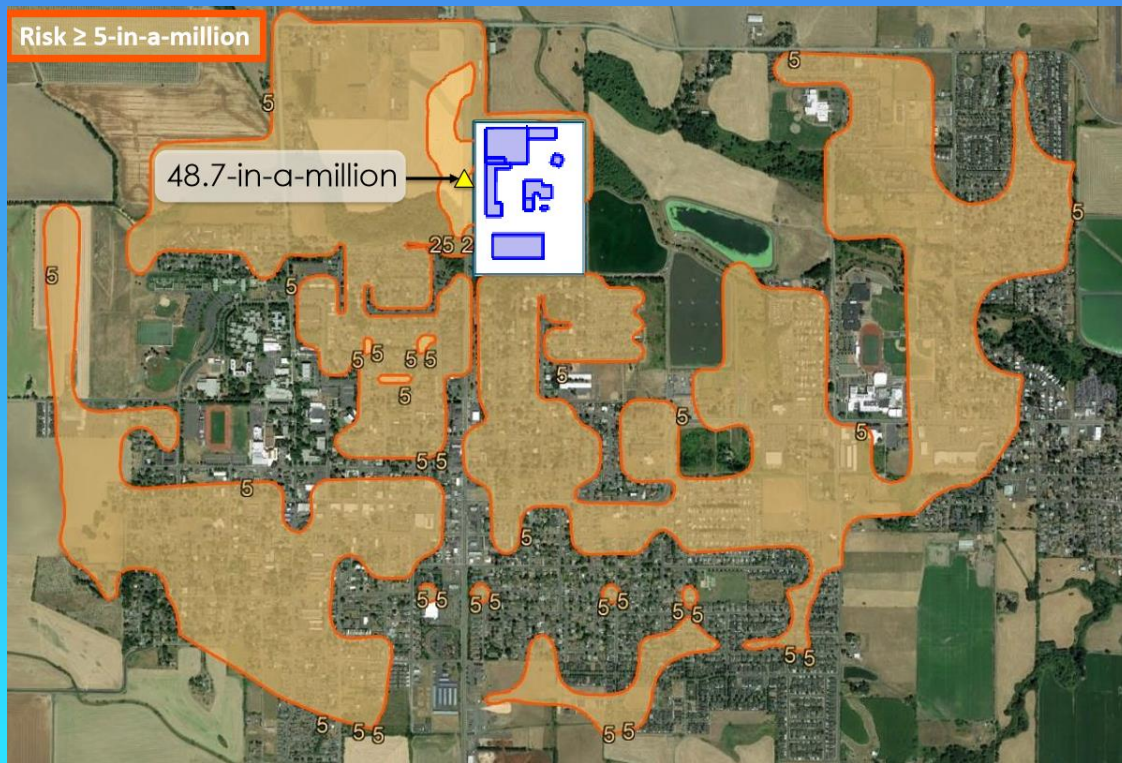
Modeling Step 3: More Math!!



2,149 receptor points
requiring data
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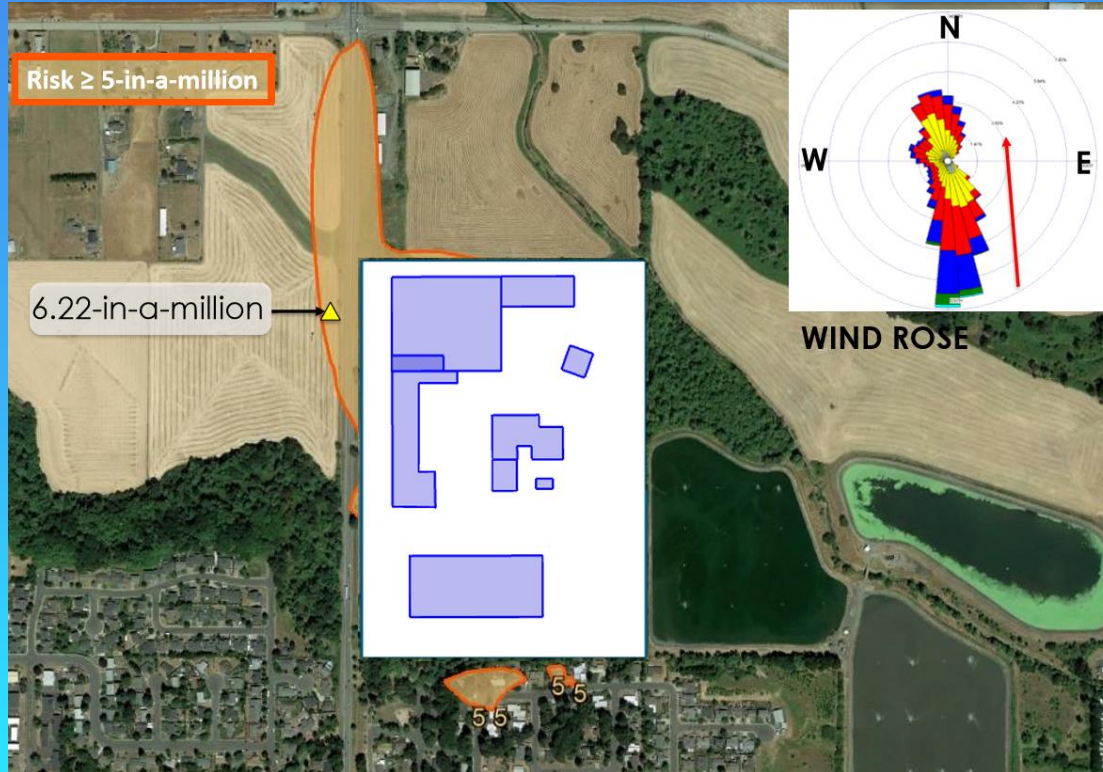
Each point on grid used
for further mathematical
modeling.

Modeling Step 4: Yet More Math!!



2,149 receptor points in conjunction with **terrain modeling** used to mathematically model isopleths.

Modeling Step 5: Math with Met Data!!



[stage whisper]:

What's the uncertainty
of the **meteorological**
data?

Collect Sample



Laboratory Analysis



Data Reduction and Modeling

Sources of Uncertainty

Sampling[‡]

- Length of sample run(s)
- Number of runs
- Scale and quality of instrument calibrations
- Sample loss due to leaks
- Analyzer drift
- Interfering gases
- Accuracy of O₂/CO₂ measurements
- Measurements of pressure and temperature
- Pitot specifications
- Non-uniform distribution of pollutants in stack
- Experience/skill of testers
- Flow measurements
- Quality of reagents
- Quality of gas standards
- Sample bottle type and cleanliness
- Glassware preparation
- Environmental conditions
- Moisture
- Field balances and other standards
- Reference balance and other reference standards
- Thermocouples
- Sampling location
- Number of points/port used
- Meter volume
- Filter efficiency and material
- Size and alignment of the nozzle during sampling
- Sample hold time and handling
- Instrument precision and accuracy
- Human error

Analysis (Laboratory)

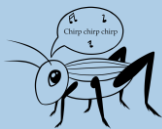
- Preparation
- Analysis

Data Modeling

- Meteorological data
- Distance/Height measurements
- Terrain maps
- Compounding uncertainties during data reduction

Sources of Uncertainty Contained in Final Reported Detection Limit

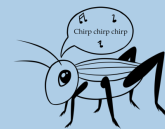
Sampling



Analysis (Laboratory)

- Preparation
- Analysis

Data Modeling



Sources of Uncertainty

Sampling[‡]

- Length of sample run(s)
- Number of runs
- Scale and quality of instrument calibrations
- Sample loss due to leaks
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- Reference balance and other reference standards
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- Sampling location
- Number of points/port used
- Meter volume
- Filter efficiency and material
- Size and alignment of the nozzle during sampling
- Sample hold time and handling
- Instrument precision and accuracy
- Human error

Analysis (Laboratory)

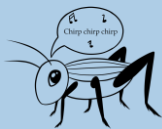
- Preparation
- Analysis

Data Modeling

- Meteorological data
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Sources of Uncertainty Contained in Final Reported Detection Limit

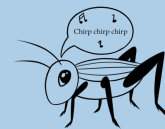
Sampling



Analysis (Laboratory)

- Preparation
- Analysis

Data Modeling



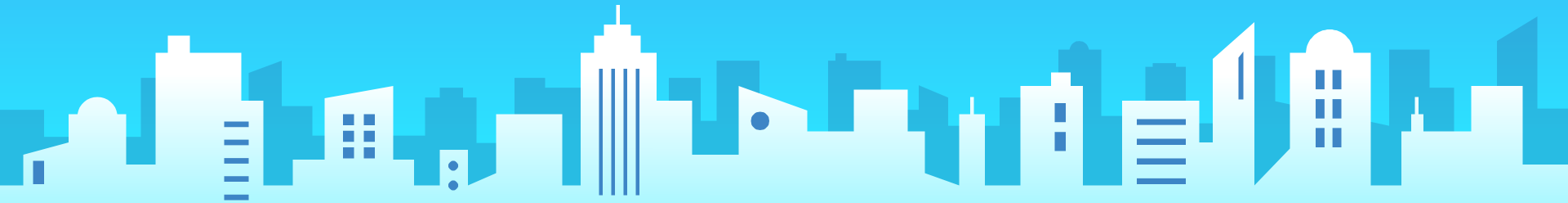
BUT WAIT! THERE'S MORE!!

Do my detectable results indicate a human health hazard?

Depending on sample collection and laboratory preparation technique, results may indicate total analyte, not bio-available analyte.

Human health hazard levels often determined by World Health Organization (WHO) – what is their uncertainty?

Is it possible to address this issue using current Source Test Methods?
(hint: probably not, at least not with current technology)



Pop Quiz

Q: Where does most of the uncertainty lie in Source Sampling final results?

A: Not with the lab.

Q: At what stage in the process are the detection limits determined that are used in reporting final results?

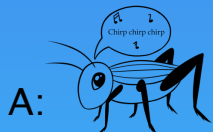
A: At the lab.

Q: Is ignoring the uncertainty from field sampling and data modeling going to bias the “detection limit” high or low?

A: Low – if other uncertainties were taken into account to create a true Method Detection Limit, the reported detection limit would be higher.

Conclusion (Points to Ponder)

Q: How does this low bias in Method Detection Limit affect the regulatory decision making process?



Q: Is it reasonably possible to take into consideration *all* uncertainty contributions in a Source Sampling Method?

A: Yes. Some aspects of uncertainty that are currently not considered during field testing or modeling could be considered.

AND

A: No. Technology would need to change to include all contributions.

How Certain Are You?



Hint: one of these is not being sampled, and at least one is using an incorrectly sized probe.[†]

[†]With thanks to Bill Guyton, ERM

QUESTIONS?

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