

Suggested QC Practices for On-Line Analysis

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Why do we need continuous monitoring?

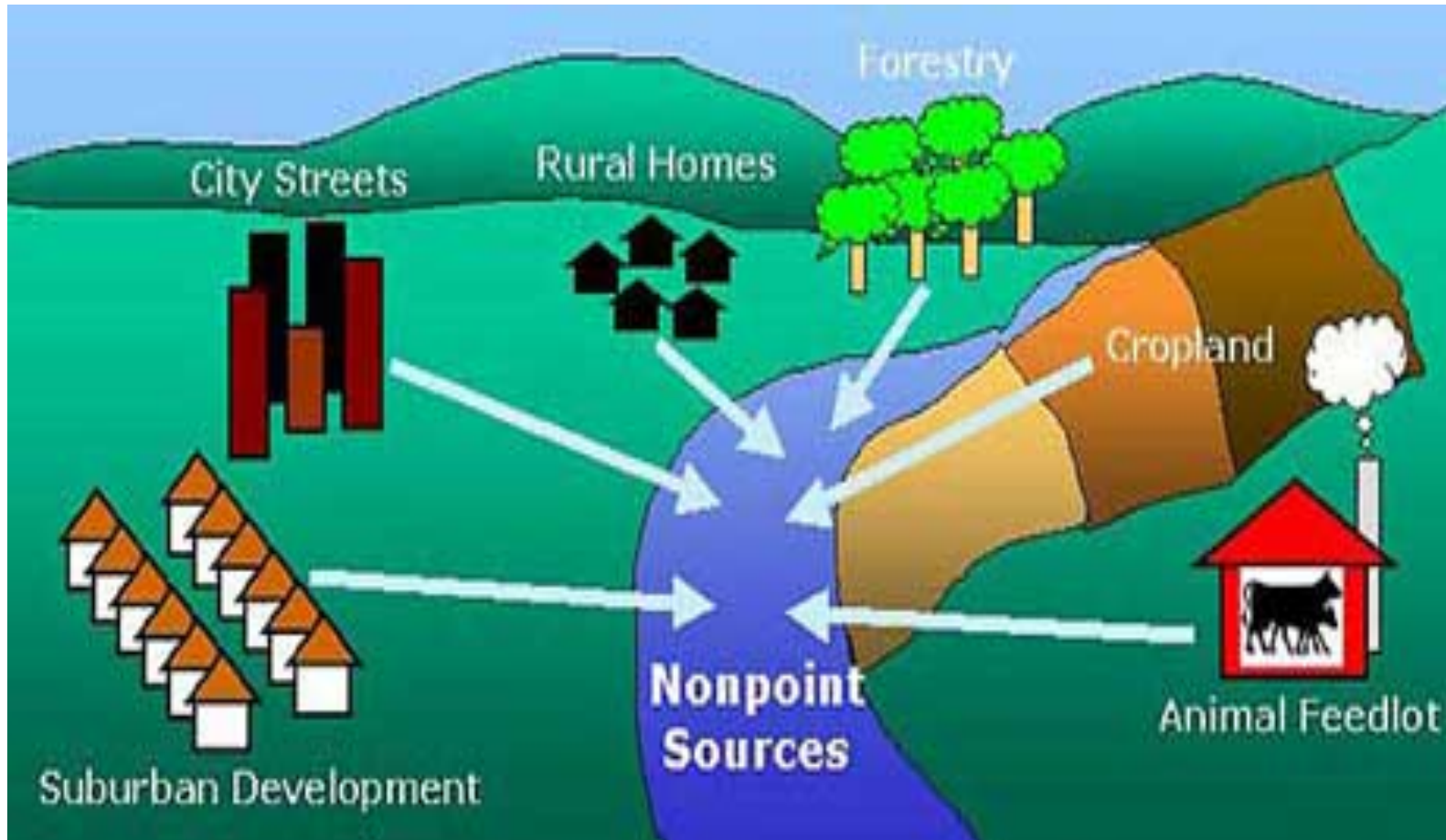


**Laboratory sampling
is only a “snapshot”
of the sample on a
good day**

Sampling at the “pipe” does not sample storm water or irrigation water



Sources of pollution may be outside “the pipe”



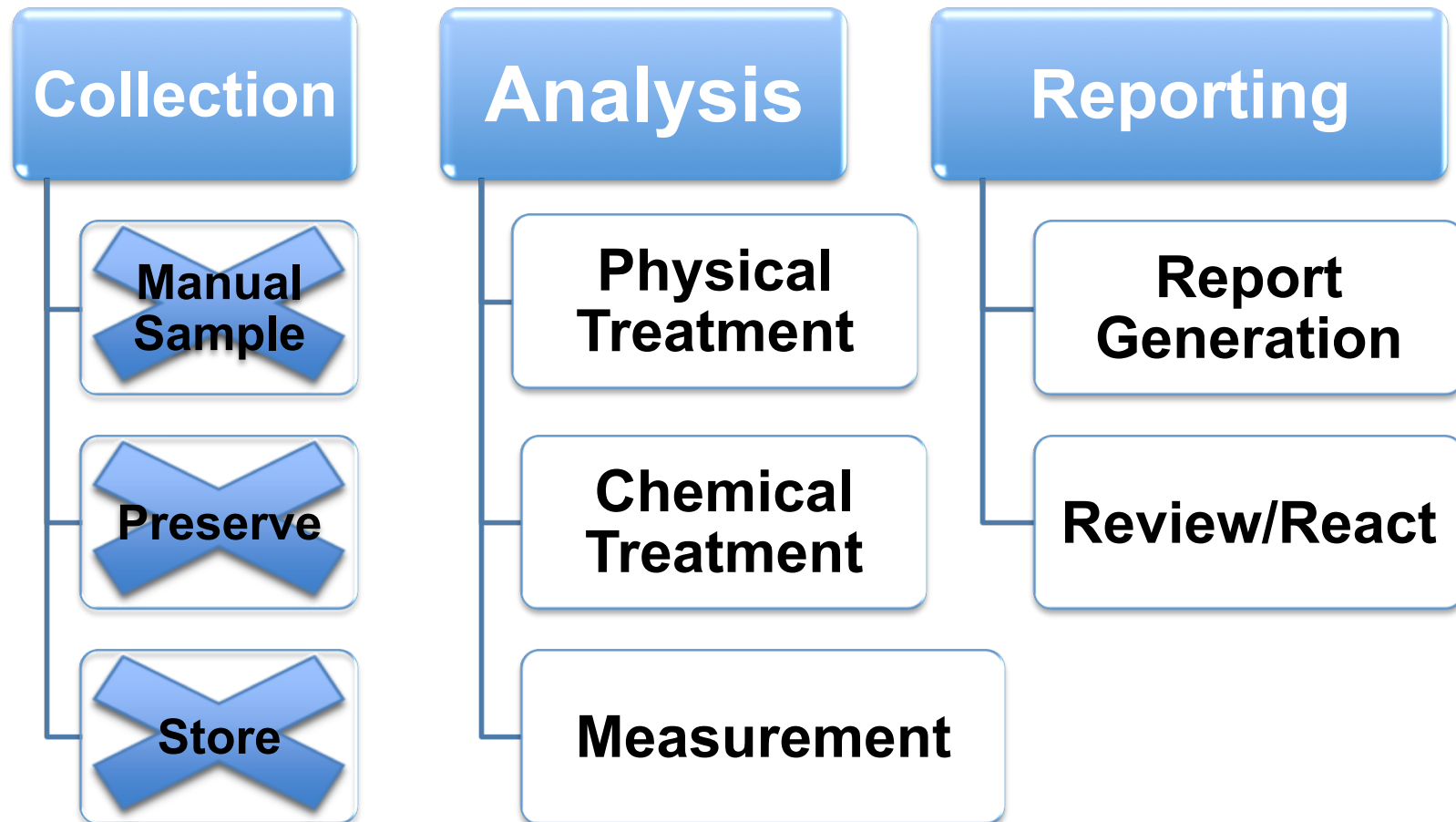
Rarely do we sample when rivers are flooding



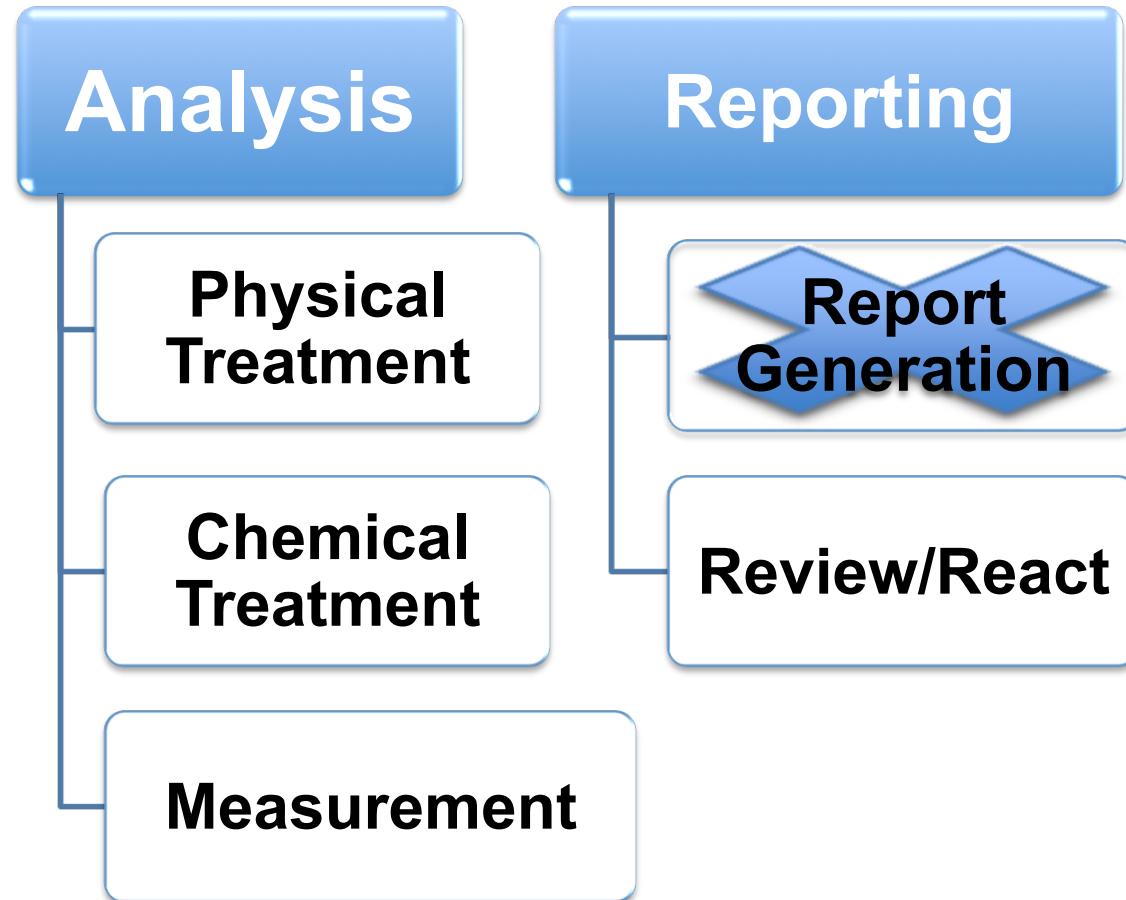
But there are no USEPA approved methods for on-line analysis!

- **EPA approves methods**
- **Methods include:**
 - **Sampling**
 - **Preservation**
 - **Calibration Verification**
 - **Duplicates**
 - **Spikes**
 - **LCS**

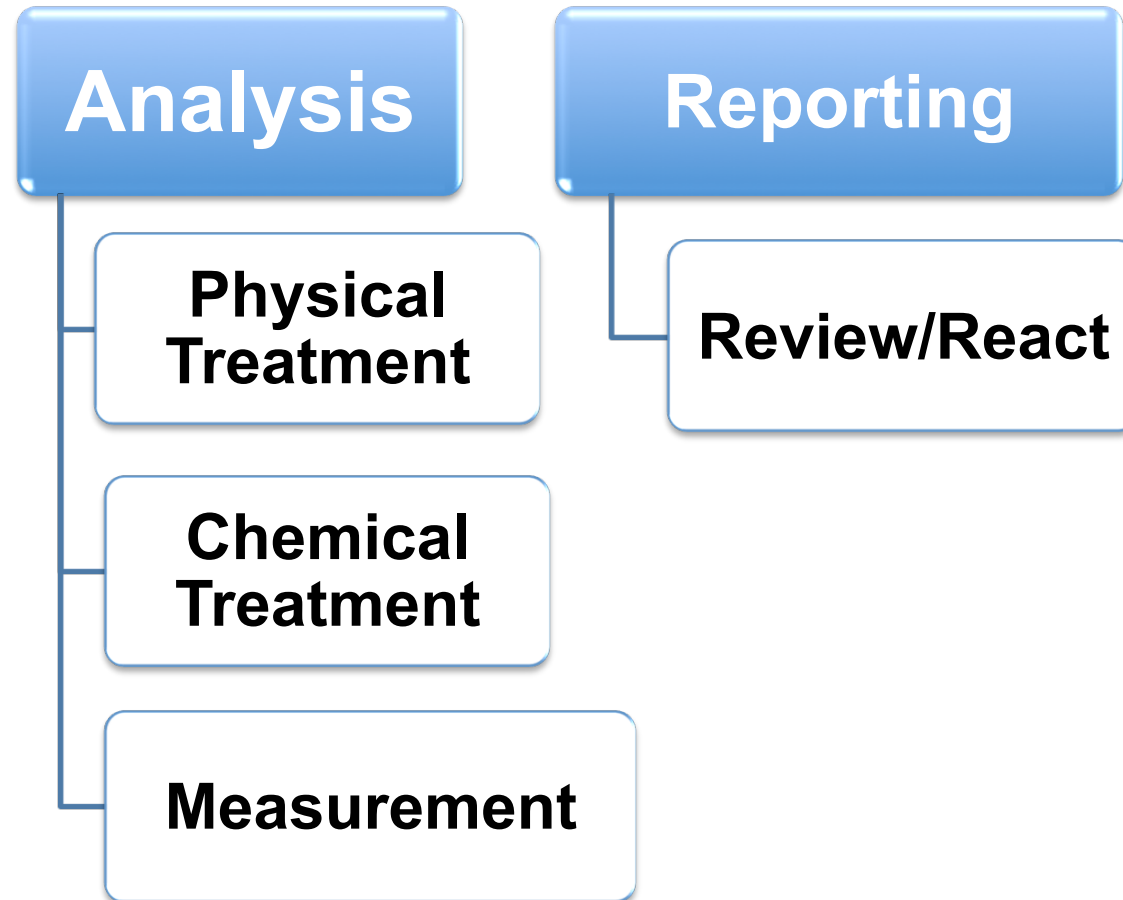
**Can get around sampling and preservation
since analysis is immediate**



**There is no generation of a laboratory report,
no batch QC**



A significant portion of “normal” is eliminated



But a method includes “batch” QC steps that must be done

- **Calibration and Calibration Verification**
- **Blanks**
- **Duplicates**
- **Matrix Spikes**
- **Analysis of Control Samples**



Calibration of on-line analyzers can be automated

- **Methods should ensure that analyzers automatically recalibrate**
- **Use multiple points if necessary**
- **Or, Operators should periodically recalibrate**

Calibration verification should be automated

- **Methods should ensure that analyzers automatically verify the calibration in a fixed interval**
- **Failed CCV should repeat calibration or flag**

Analyzer should run “blanks” if target analyte varies

- **Process control**  **blanks**
- **Ambient water**  **blanks**

**The concept of “duplicates” does not apply,
however precision should be evaluated**

- **The on-line method should require a way to collect repeatability data**
- **Stick the sampling straw in a beaker**

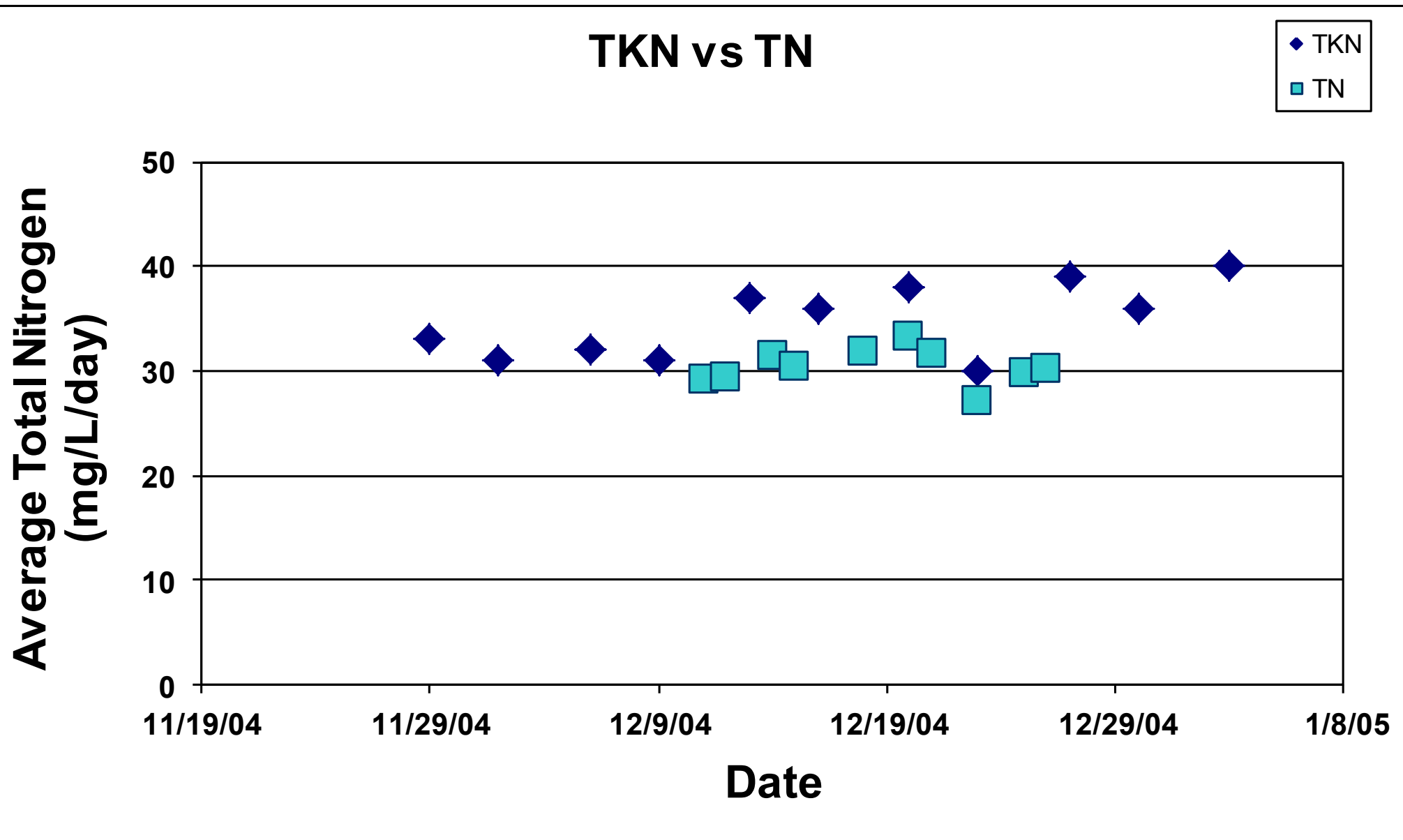
**The concept of “spikes” does not apply,
however recovery should be evaluated**

- **The on-line method should require a way to determine recovery in the matrix**
- **Recovery determined on commissioning**

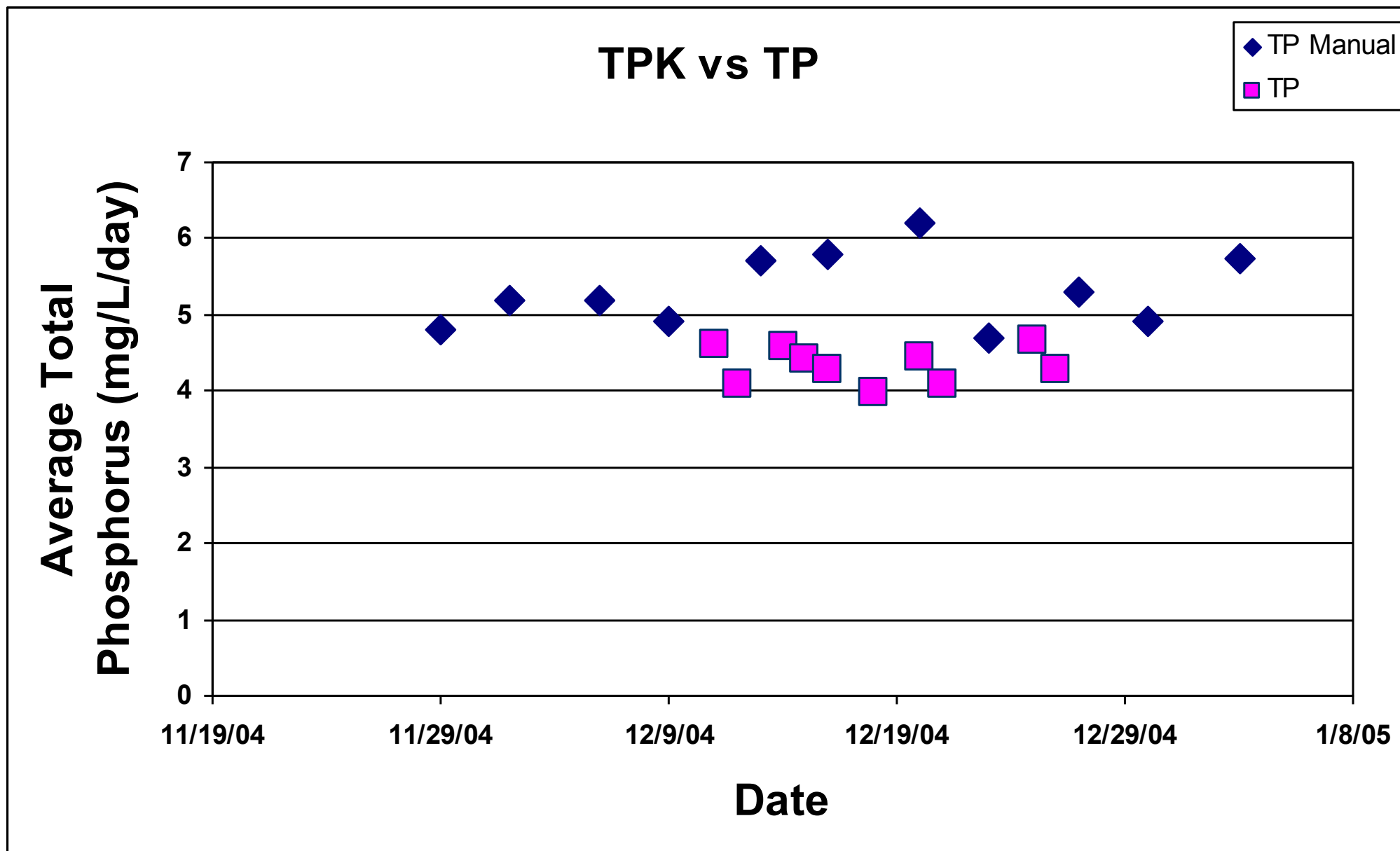
The concept of “LCS” does not apply, however precision and recovery should be evaluated

- **The on-line method should periodically evaluate an LCS**
- **LCS defined as comparison to approved method**

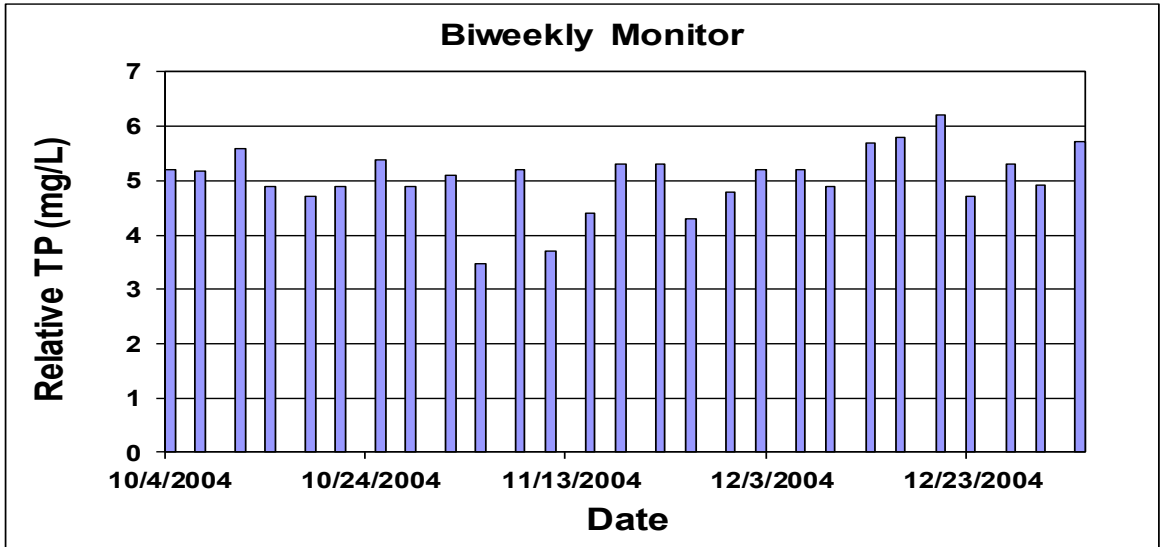
Example of TN (on-line) compared to TKN (approved)



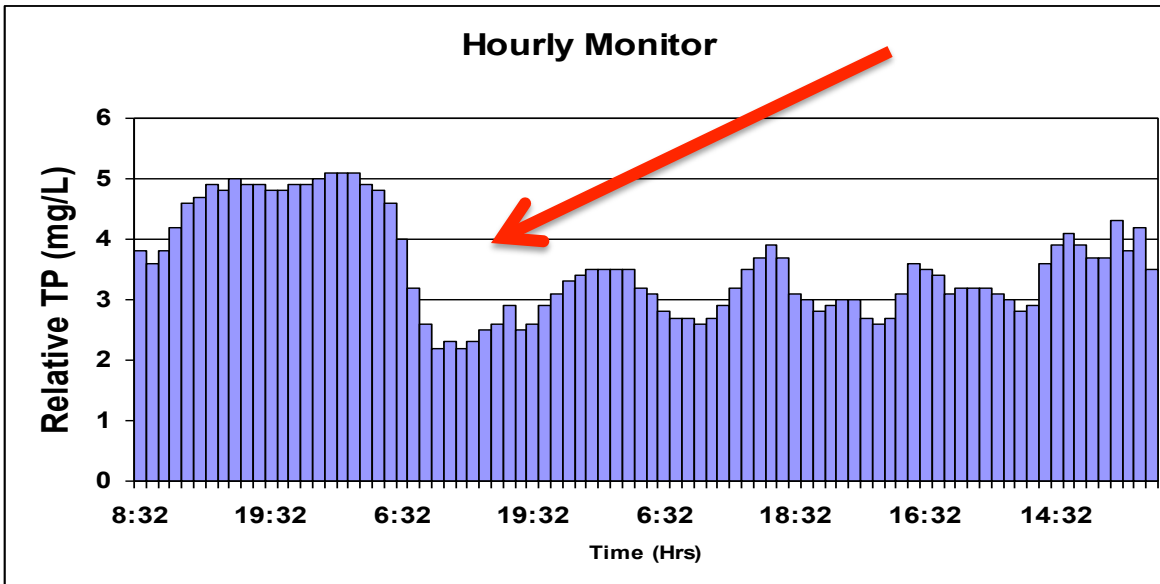
Example of TKP (approved) compared to TP (online)



On-line monitoring allows you to see “upsets”



Laboratory



On-line

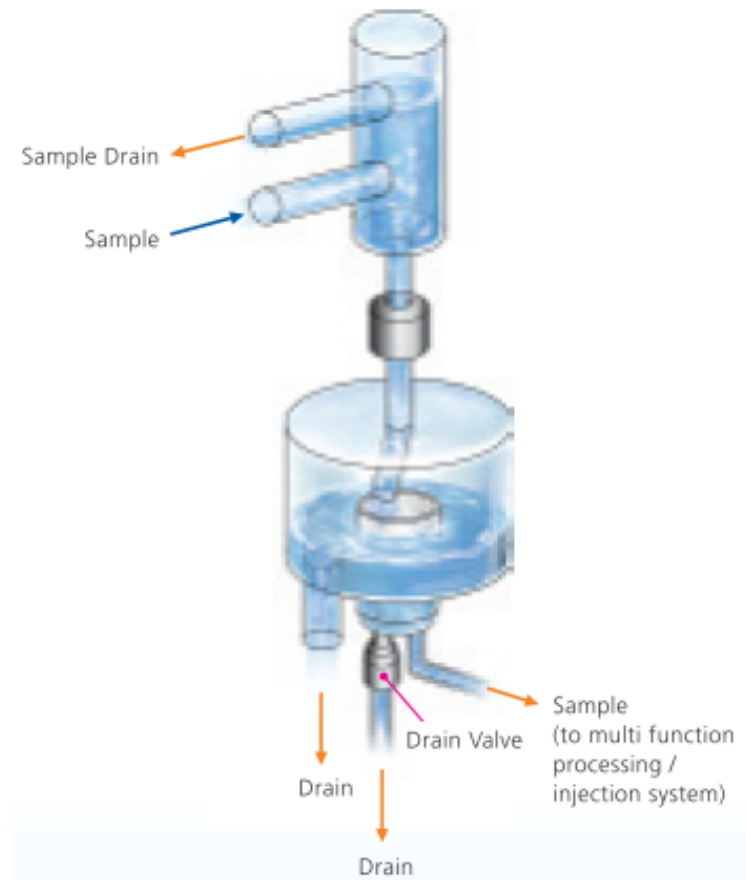
Any new “method” for continuous on-line monitoring should:

- **Obtain equivalent results to approved laboratory methods**
- **Be “rugged”**
- **Use few reagents**
- **Have a sufficient range**
- **Sample/”handle” a complex matrix**
- **Calibrate with multiple points**
- **Require periodic check samples compared to lab**

The on-line analyzer must be rugged enough for field/plant use

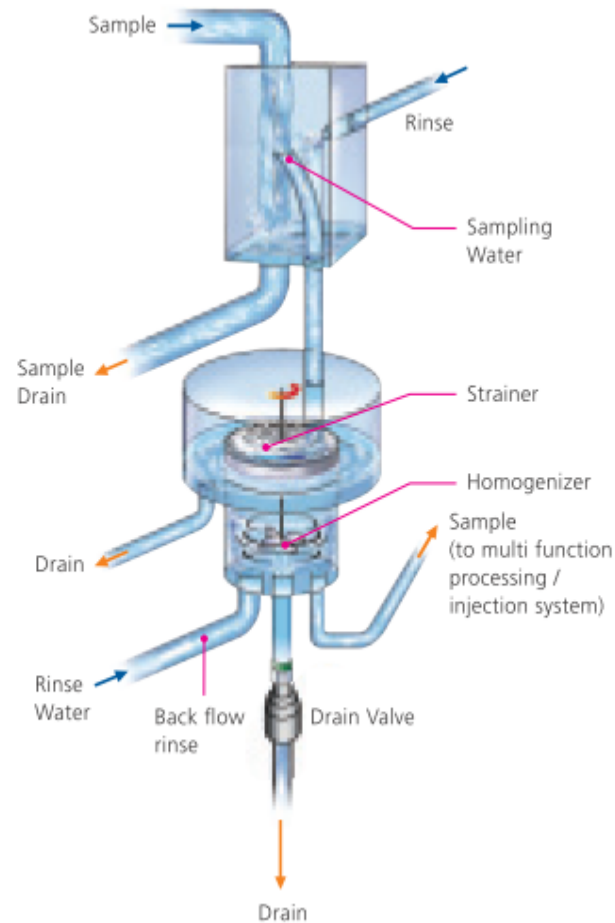


The on-line analyzer should be capable of sampling multiple streams, or cost effective



Multi-Stream Sample Switching Unit

The on-line analyzer should be capable of handling TSS if necessary



Multiple Stream TSS Unit

Benefits of On-Line Monitoring

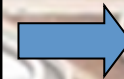
Automatic

- Sampling
- Pretreatment
- Digestion
- Analysis
- Report



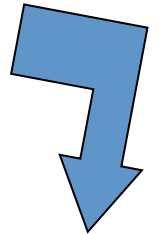
Flexibility of Sampling

- Hourly
- Daily
- Weekly
- Monthly



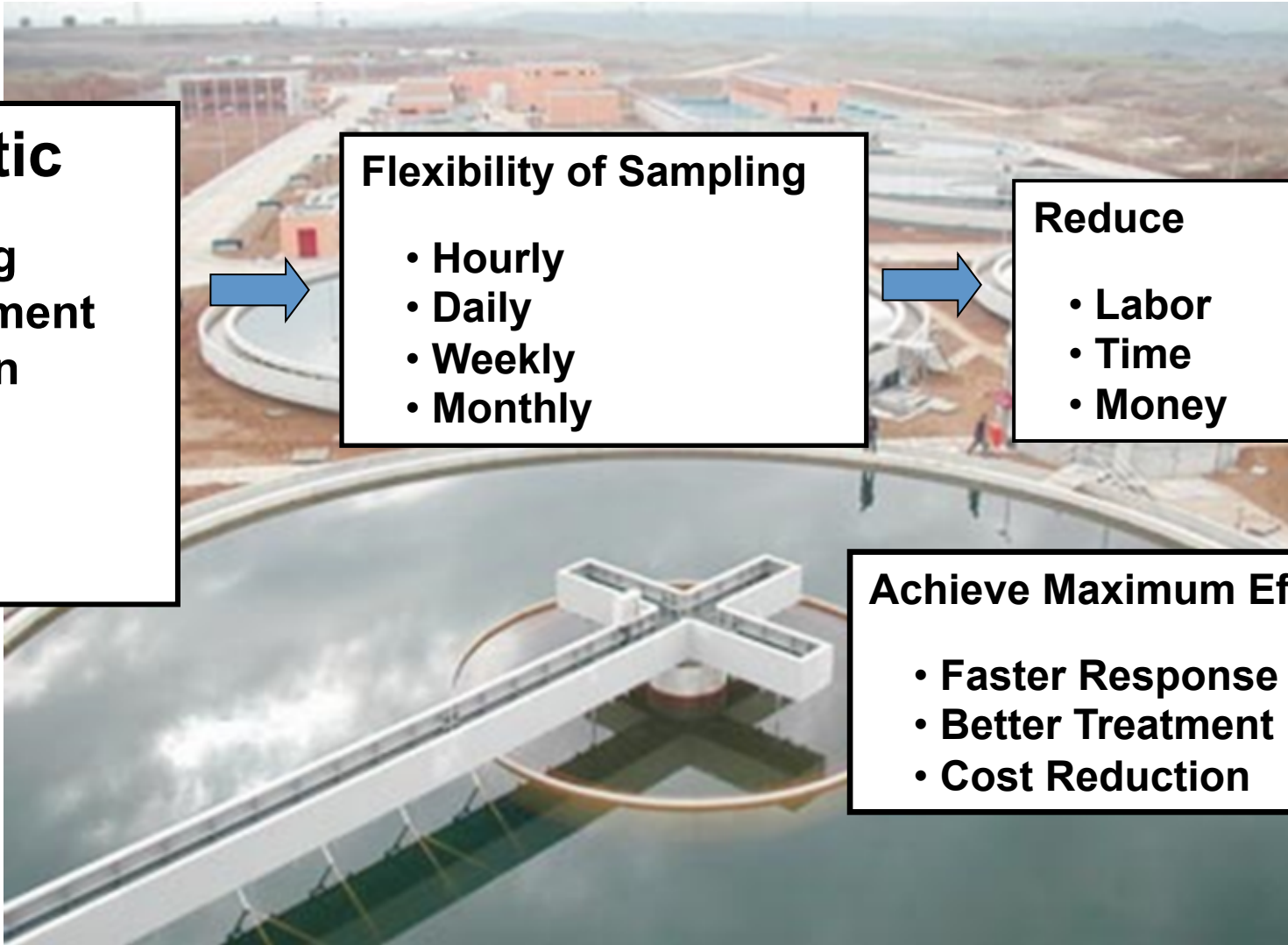
Reduce

- Labor
- Time
- Money



Achieve Maximum Efficiency

- Faster Response
- Better Treatment
- Cost Reduction



On-line analyzer methods are needed if data is to be reported for compliance

- **Some QC in existing methods does not apply**
- **New methods can be written that still provide sufficient QC**
- **On-line results should be compared to lab results**

Thank You!

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For more information contact

