



VOC Emission Testing

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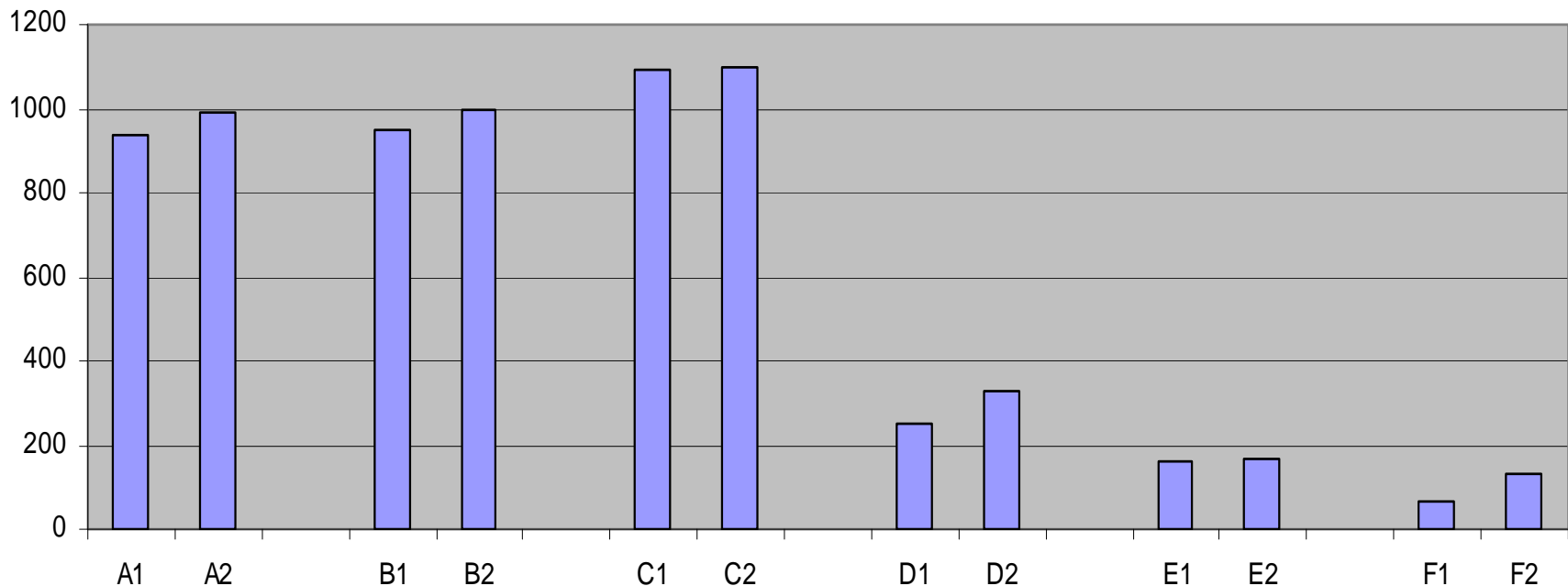
Some quality issues:

Reproducibility from lab to lab

Reinhard Oppl, Eurofins Product Testing

Within lab reproducibility

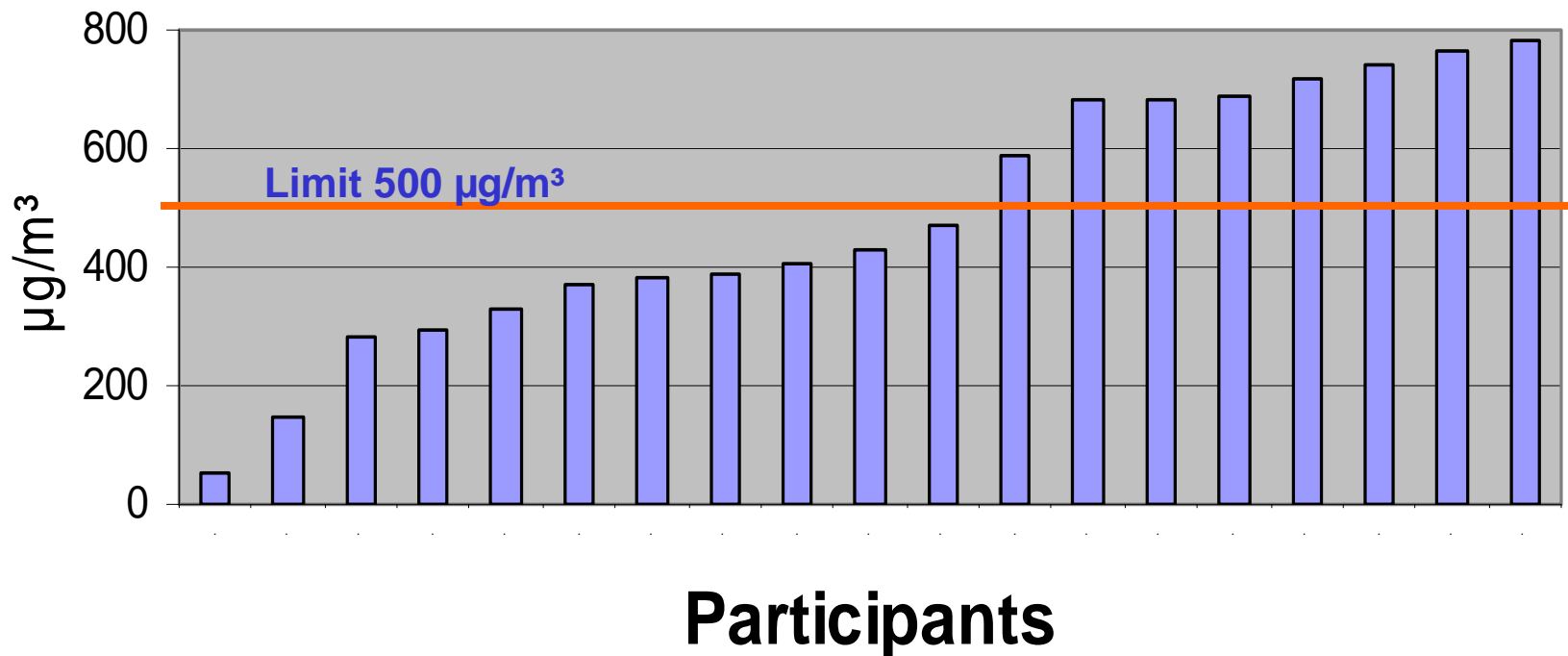
Typical TVOC reproducibility within one laboratory:
6 flooring samples in 2 test chambers each ($\mu\text{g}/\text{m}^3$)



A - F: Different samples. 1, 2: Two test specimens taken from same sample.

Reproducibility – 20 labs

TVOC 2nd GEV Round Robin Test - 1 test adhesive TVOC GEV (Results of Participants)

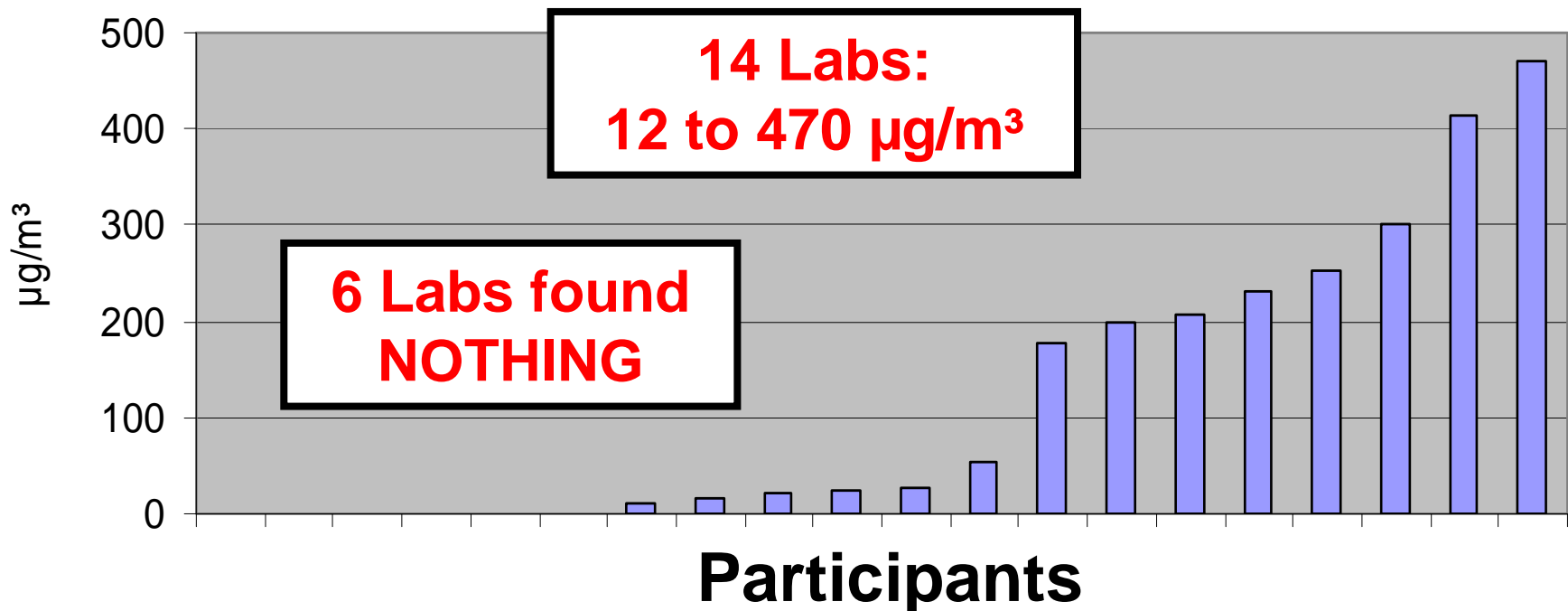


1 lab did not report a TVOC

Reproducibility – Single substances

2. GEV Round Robin Test

Ethylene glycol: 20 labs testing ONE adhesive



Status

- **Reproducibility of VOC test results:**
 - Variation $\pm 20\%$ (RSD) should be guaranteed within one lab (or better - but limited by material inhomogeneity)
 - But: Low concentration = higher uncertainty (weaker detector signal)
 - Variation $\pm 50\%$ between different labs (RSD)
Some improvement should be possible ($\Rightarrow \pm 20\text{-}30\%$ for homogeneous material) - but this is **always limited by material inhomogeneity**
- **Other than VOC**
 - SVOC, VVOC, reactive compounds (e.g. formaldehyde) cannot be measured in a reliable manner with the VOC method, these require different analytical methods
- **Air concentration below 0.005 mg/m^3**
(for some compounds even already below 0.02 mg/m^3)
are not significant at all in many cases

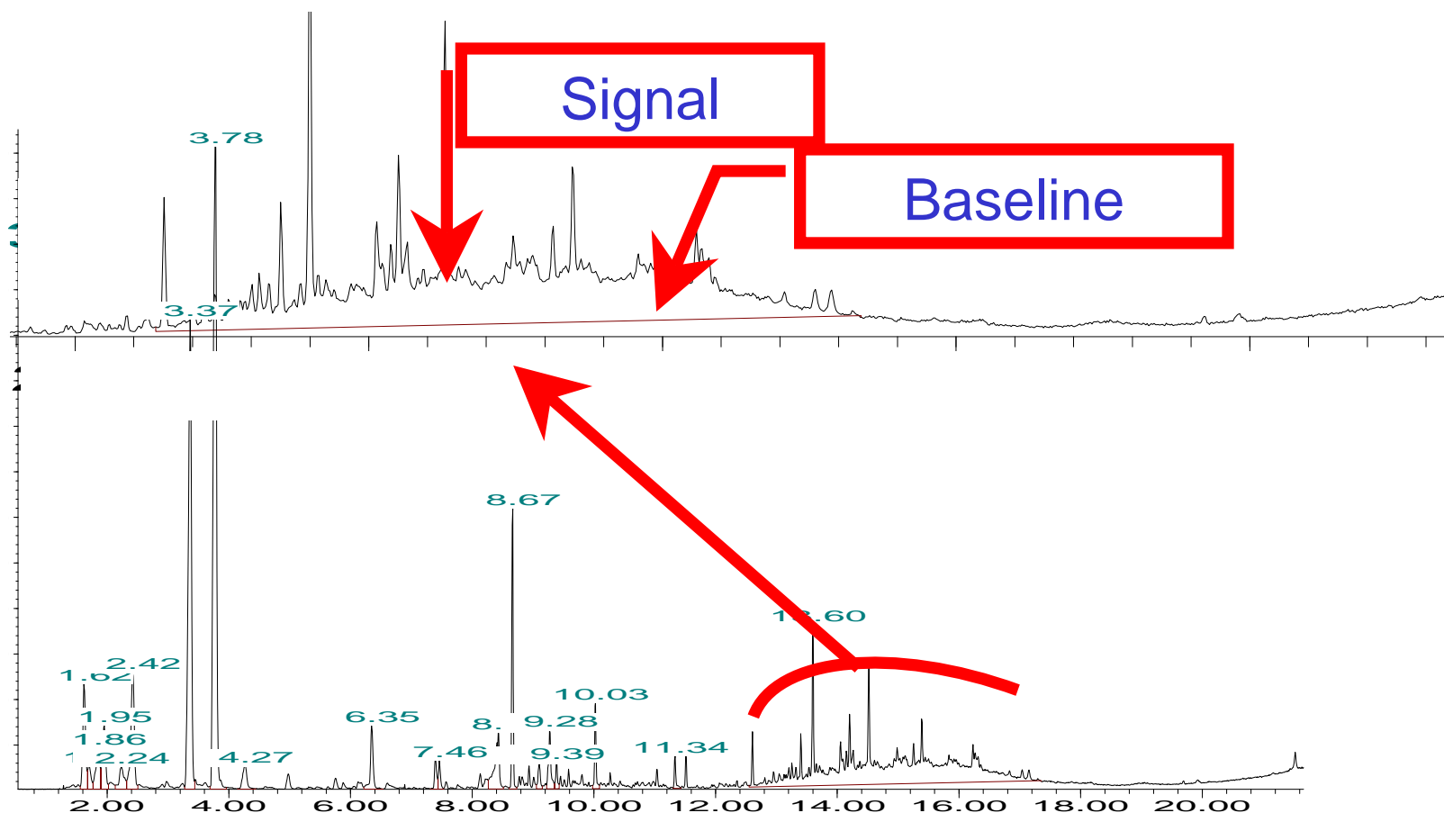
Starting point

- **Tested product**
 - **Inhomogeneous emissions over surface or over batches**
 - **This cannot be changed**
 - can only partly be levelled out by using larger chamber in case of known higher inhomogeneity over surface
- **Gas chromatographic analysis** (see slide 9)
 - **Time corresponds with identity, to be confirmed by MS**
 - **Height corresponds with mass of substance**
 - **Different VOC give different signal per mg ("response factor RF")**
 - Day in - day out calibration with toluene, high sequence
 - Calibration of other VOC than toluene: With its "relative response factor RRF" - relative to toluene - e.g. RRF 1.25 means toluene result is corrected by division with 1.25
 - The RRF values are pretty stable and must NOT be checked daily, can even be re-applied later on stored old data

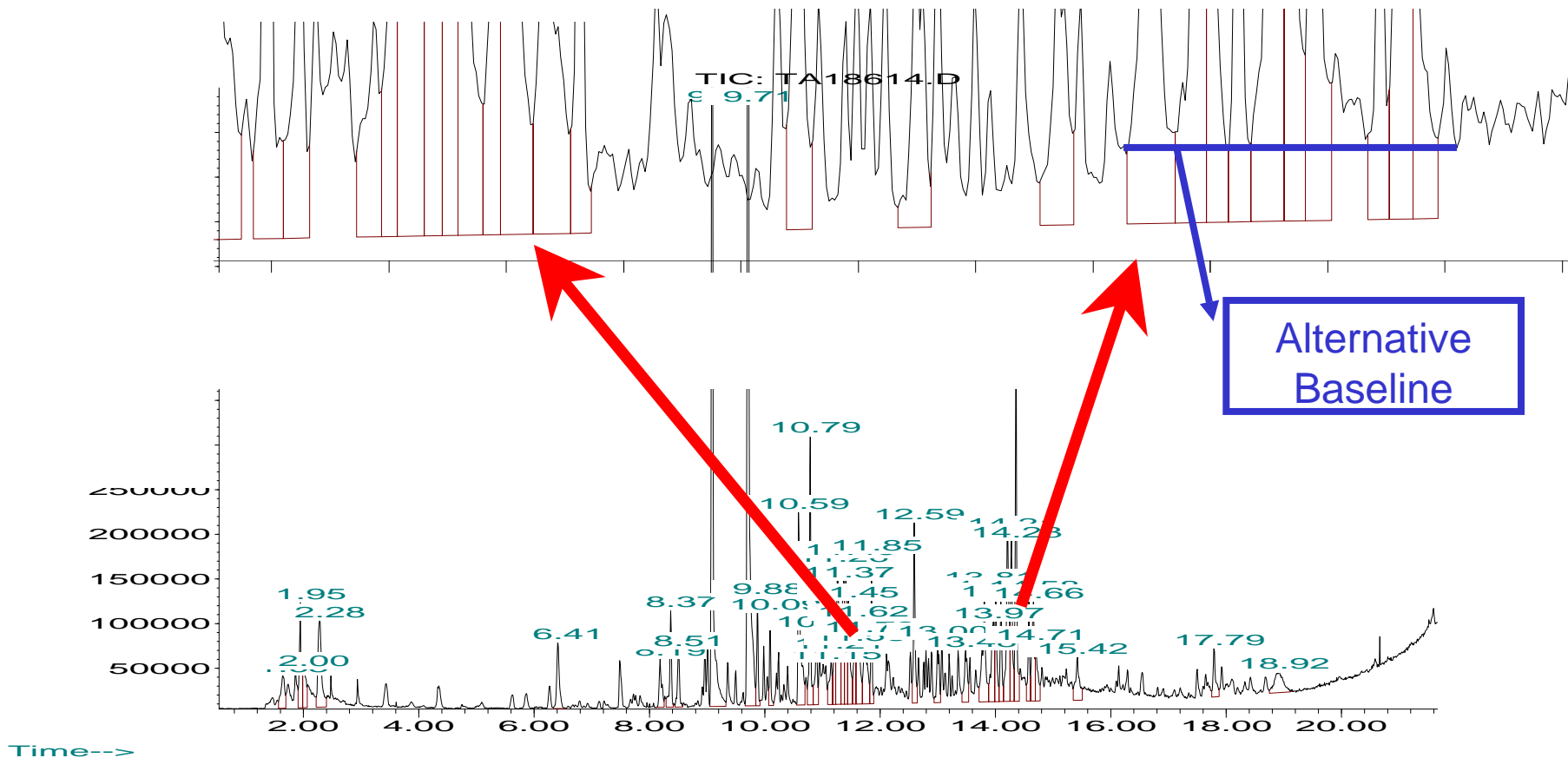
Key issues I (GC analysis)

- **Different separation of VOC mixture gives different chromatograms**
 - ISO 16000-6 allows use of different columns
 - Should be restricted to 100% non-polar column (see VOC definition)
- **Different detectors give different quantitative signal as toluene RF**
 - ISO 16000-6 allows use of different detectors (MS or FID)
 - Should be restricted to one detector only (MS)
- **Different calibration factors give different results**
 - ISO 16000-6 requires 2/3 of signal to be identified
=> differences per lab and its identification power. Better is:
 - Use only toluene equivalent, or
 - Use specific calibration for a list of target VOC,
and toluene equivalent for all other substances
- **Different peak baselines give different quantification**
 - Harmonise interpretation procedures, see next slides

Identification and Quantification – different interpretation possible



Quantification in complex mixture – different interpretation possible



Key issues II (air sampling)

- **Air sampling from test chamber: 2x2 samples per result**
 - Always apply and analyse backup sampling after front sampling, and report sum of both sampling tubes
 - for identifying any breakthrough in case of high emissions
 - Otherwise: High risk of erroneous results
 - Discard whole test if >25% breakthrough
 - Backup tube may be skipped if low emissions are expected
 - Always apply parallel double sampling, and report average of both samplings
 - for identifying any error in sampling,
 - Otherwise: High risk of erroneous results
 - Discard whole test if >20% difference
 - 2 samplings after each other acceptable if only 1 sampling port is available



Key issues III (more air sampling)

- **Selection of adsorbent for VOC analysis**
 - **Different sorbent gives different**
 - adsorption capacity
 - desorption efficiency
 - selectivity during sampling and analysis
 - **Use only one sorbent: Tenax TA for VOC (as defined in ISO 16000-6)**
 - Plus: Harmonisation between labs
 - Minus: Not so good for VVOC, SVOC
- **Sampling time and duration**
 - When emissions are still in decay, slightly different sampling intervals may determine result
 - Define starting date / hour, then allow small intervals only



**Testing with high reliability
will lead to less questions and doubts**

=>

**saves money compared to
uncertain tests**

even if

**high quality tests may be
more expensive**

Validation of Test Method and of Quality Criteria for Test Houses are essential

Further reading:

R.O ppl: Reliability of emission testing,
Gefahrstoffe Reinhaltung der Luft 68(2008)3 pp 83-86

Free download:

[http://product-testing.eurofins.com/
testing/chemical-safety/emission-into-indoor-air.aspx](http://product-testing.eurofins.com/testing/chemical-safety/emission-into-indoor-air.aspx)