



ANALYTICAL TECHNIQUES, USED IN EXTRACTABLES TESTING

PDA TRAINING COURSE EXTRACTABLES – LEACHABLES Rome 01 – 02 March, 2018

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CHALLENGES IN E/L TESTING



Challenges in E/L-Testing

Diversity of not-API Related Compounds in E/L research is Tremendous!!

Broad spectrum of:

- Types of Containers
- Types of Materials used in the Manufacture of Containers
- Number of Suppliers per Material
- Number of Grades (per supplier) for each type of Material
- Type of Sterilization (impact on material impurity profile)

PDA Challenges in E/L-Testing

(Non Limitative) List of types of Pharmaceutical Containers

INHALATION

- o Metered Dose Inhaler Components
 - e.g.:
 - Gaskets
 - Stem
 - Body
 - Metering Chamber
 - Protection Ring
 - Actuator
 - Cannister
- o Dry Powder Inhaler Components
- Nasal Spray Systems
- Nasal Dropper Systems
- 0 ...

PARENTERAL

- o Bottles
- o Vials
- o (Pre-Filled) Syringes
- o Cartridges
- o (Rubber) Stoppers
- Rubber Plungers
- o Needle Shields
- \circ Tip Caps
- o I.V. Bags
- o Administration Sets
- 0 ...

OPHTHALMIC

- Eye Dropper Systems
- o Tubes
- 0 ...

DERMAL/TOPICAL

- o Spray Systems
- \circ Tube systems
- 0 ...

SINGLE USE SYSTEMS

- o (Multilayer) Bags
- o Tubings
- o Connectors
- o Ports
- Filters (+ Housing)
- Chromatographic Columns
- o Lyo trays
- 0 ...

SECONDARY PACKAGING

- o Labels
- o Adhesive/Glue (e.g. on labels)
- o Ink
- o Overwrap foils
- o Blisters
- o Cardboard packaging
- 0...

DA Challenges in E/L-Testing

Pharmaceutical Containers can be made of different Materials

- Low Density Polyethylene
- High Density Polyethylene
- Polypropylene
- \circ Rubbers
- o Butyl Rubbers
- Chlorobutyl Rubbers
- \circ Bromobutyl Rubbers
- EPDM Rubbers
- \circ Isoprene Rubbers
- o Nitrile Rubbers
- Latex Rubbers
- \circ Other Rubbers
- o Multi-layer Films and Foils
- Polyurethane (PU)
- Ethylvinyl Acetate (EVA)
- Ethylvinyl Alcohol (EVOH)

- Polyamide (Nylon-6, Nylon-66)
- Cyclic Olefin Copolymers (COC)
- \circ Cyclic Olefin Polymers (COP)
- Polyethylene Terephthalate (PET, PETG)
- Polybutylene Terephthalate (PBT)
- Polyacetal (POM)
- o Polymethylmethacrylate (PMMA)
- Acrylonitrile Butadiene Styrene (ABS)
- o Silicone
- $\circ \text{ C-Flex}$
- o Polycarbonate
- \circ Teflon
- \circ PEEK
- \circ Glass
- \circ Metals
- 0...



Challenges in E/L-Testing

Each Material has different Suppliers

EXAMPLES

Polyethylene - produced by:

- o Borealis
- o LyondellBasell
- o SABIC
- o Dupont
- o Enichem
- o INEOS
- o TOTAL
- 0 ...

Pharmaceutical Rubbers - main Global Suppliers:

- o Datwyler
- o West Pharmaceutical
- o Stelmi

Each Supplier has different Different Grades!



Each Supplier has different Different Grades

EXAMPLES

PolyEthylene - produced by:

- o Borealis: over 30 different Medical Grades
- o LyondellBasell: over 30 different Medical Grades
- o SABIC: over 30 different Medical Grades
- o Dupont: different grades
- o Enichem: different grades
- INEOS: different grades
- o TOTAL: different grades
- 0 ...

Pharmaceutical Rubbers - main Global Suppliers:

- o Datwyler: over 100 different commercial rubber formulations
- West Pharmaceutical: over 100 different commercial rubber formulations
- o Stelmi: also, a broad range of commercial rubber formulations



Challenges in E/L-Testing

Per Material, Supplier and Grade: what makes up the Impurities Profile?

- Solvent residues (e.g. of Polymerization)
- Polymer residues (e.g. Monomers, Oligomers)
- Catalysts
- Polymer/Rubber Additives
 - \circ Antioxidants
 - \circ Photostabilizers
 - o Plasticizers
 - o Lubricants
 - Acid Scavangers
 - Pigments/Colorants
 - Carifying/Nucleating Agents
 - Cross Linking Agents (Rubbers)
 - o Initiators (Rubbers)
 - Accelerators (Rubbers)
- Polymer Additive Degradation Products
- Polymer Degradation Compounds
- Adhesives
- ≻ ...



Conclusion:

- The <u>broad diversity</u> of pharma containers, materials, suppliers and grades, leads to a extremely <u>long list of potential impurities</u> (leachables), introduced into the drug product
- 2. The <u>compounds cannot be investigated with 1 analytical technique</u>. Typically, at least 3 to 5 analytical techniques will need to be combined.
- 3. Compound Identification is of high importance, therefore the <u>detection</u> <u>needs to be compound specific</u> (e.g. MS-detection)
 - Headspace GC/MS Volatile Organic Compounds
 - GC/MS Semi-Volatile Organic Compounds
 - LC/MS Non-Volatile Organic Compounds
 - ICP Metals
 - IC Anions





Conclusion:

4. For Companies / Labs, only performing E/L-testing, <u>every E/L-project</u> could turn out into a <u>high level research project</u> (with the need for high level analytical techniques) <u>because of the lack of materials knowledge</u>

 For Labs, performing E/L-studies on a routine basis, excessive analytical costs (associated with high-end analytical procedures) should be avoided in FIRST PASS testing. Toxikon: TOX-RAY development





ANALYTICAL TECHNIQUES USED FOR EXTRACTABLES TESTING





SAMPLE PREPARATION:

THE MOST IMPORTANT & THE MOST UNDERESTIMATED ACTIVITY IN THE LAB!!!

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SAMPLE PREPARATION – CHALLENGES IN TRACE ANALYSIS

- Have very experienced people in Sample Preparation
- Very Intensive Training for new staff in Sample Prep
- QC on solvents used select batches of clean solvents with suppliers
- QC on extraction equipment
- Separate glassware
- Precleaning of glassware validation of Cleaning Procedures
- Sampling of test articles how to handle Test Articles?
- WFI sample prep should be separated from solvent sample prep
- Correction for absorbed solvents?
- How to **concentrate extracts** while avoiding cross contaminations
- Storage of extracts under controlled conditions
- Holding times of extracts
- Selection of type of containers for storage of extracts
- How to keep **DEHP** out of the Lab!

PDA ANALYTICAL TECHNIQUES – SAMPLE PREP



- How to deal with human source contaminants (limonene, squalene, parabens, palmitic/stearic acid...)
- Headspace GC/MS: WFI should be completely SEPARATED
 - o Sample prep
 - Storage of sample/extract
 - Filling into storage containers
 - o Instruments
 - Holding times for HS-GC/MS are shorter!!
 - Avoid cross contamination from other solvents, regularly used in the lab (DCM, Hexane, IPA, Toluene, Chloroform...)
- o Internal standards
 - Holding times of Internal Standards
 - Syringes: should be calibrated at least yearly
 - Have a cleaning procedure for syringes
 - Compatibility of Internal Standards with solvents





EXTRACTABLE STUDIES

IDENTIFICATION

IDENTIFICATION

IDENTIFICATION





INCREASE THE KNOWLEDGE ABOUT THE COMPOSITION OF THE POLYMER



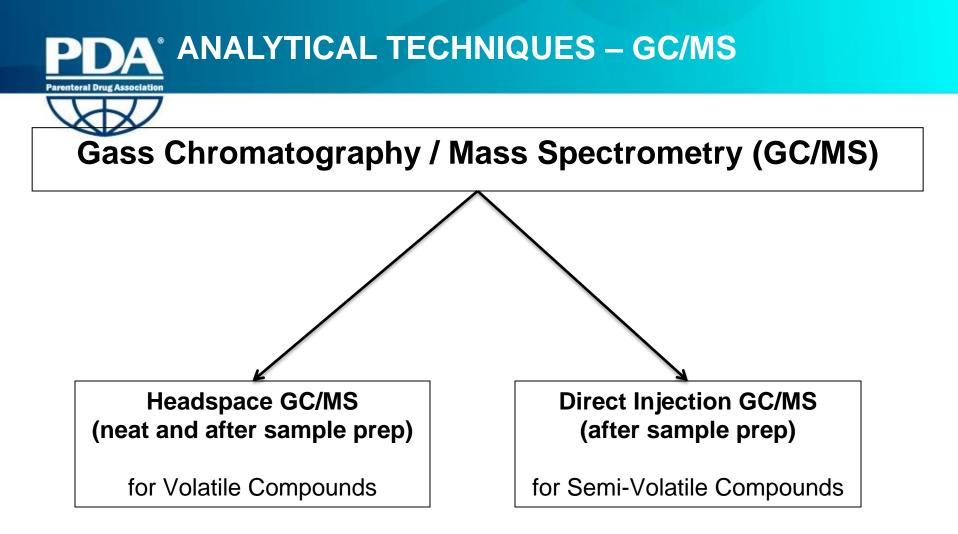
- ADDS TO INFORMATION PROVIDED BY RAW MATERIAL SUPPLIERS OR C/C MANUFACTURERS
- EXTRACTABLES LIST: FOCUS FOR LEACHABLE STUDY
- IN SOME CASES: QUANTITATIVE EXTRACTABLES STUDIES (e.g. inhalation)



EXTRACTABLE STUDIES

A **Broad Identification** in "First Pass" Extractable Studies Requires:

- 1. A Compound Specific Detector: Mass Spectrometry
- 2. A **Database** to allow Identification based upon Mass Spectra
 - Commercial Databases for GC/MS: NIST, WILEY
 - Self-Developed Databases (e.g. **TOX-RAY**)
 - **PROBLEM for LC/MS**: no Commercial Databases Available!





However, the GC/MS part of the Instrumentation is the same for the two techniques!!



PDA[®] ANALYTICAL TECHNIQUES – GC/MS

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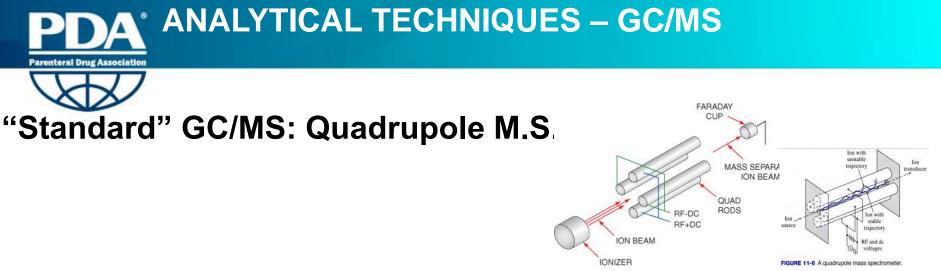
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"Standard" GC/MS: Quadrupole M.S.

Gas Chromatography: Separation of Organic Molecules based on:

- Polarity Interaction/Affinity with the Stationary Phase
- Boiling Point GC-Oven temperature
- Film Thickness of the Chromatographic Cap Ilary Column
 - \circ Volatile Compounds: high film thickness (>1 μ m)
 - \circ Semi-Volatile Compounds: low film thickness (≤0.25 μm)
- Length of the Chromatographic Capillary Column
 - \circ Volatile Compounds: 30 m to 60 m
 - o Semi-Volatile Compounds: 30 m
- Polar Organic Compounds may need more specific conditions
 - o Acids, Amines, Alcohols....





General Sequence of Things in a Mass Spectrometer (GC):

- High Vacuum
- o Convert Molecules to Ions (Tungsten Filament)
- A Moving Ion (= charge) in a Magnetic Field gets deflected
- Only the right "m/z" can reach the detector and give a (charge) signal
- $\circ~$ The charge signal is "strengthened" by a photomultiplier
- The Mass Filter (e.g. Quadrupole) scans a predefined mass range in milliseconds!
- This way, a complete mass spectrum can be obtained in a few milliseconds!



Standard GC/MS: Quadrupole M.S.

- A GC/MS "Mass Spectrometer" is <u>Standardized</u>:
 - 1. Quadrupole (or Ion Trap)
 - 2. Ionisation: Electron Impact Ionisation of 70 eV
 - 3. Gives Reproducible Mass Fragmentation:

Reproducible Mass Spectrum

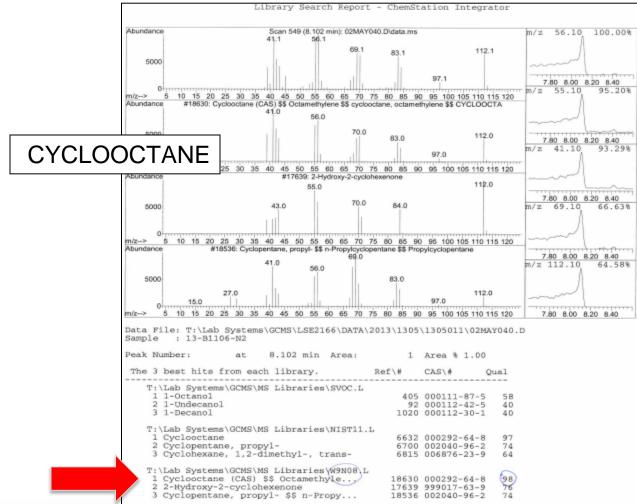
- Mass Spectrum can be compared to commercially available Databases, such as NIST or WILEY – or self-developed MS-Databases (eg TOX-RAY)
- 5. Can lead to Identification of Compound

PDA ANALYTICAL TECHNIQUES – GC/MS



Standard GC/MS: Quadrupole M.S.

Example of FIT of an UNKNOWN MS with NIST/WILEY





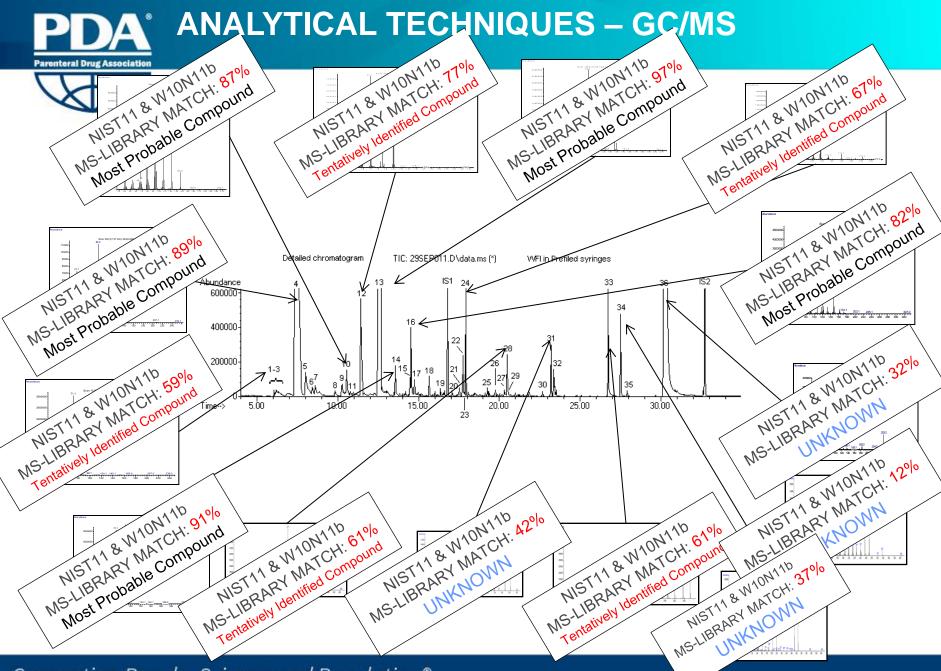


"Standard" GC/MS: Quadrupole M.S.

WHAT IS "SCREENING"?

- Trying to identify every single peak in a chromatogram
- Above a certain threshold
 - either Analytical (reporting threshold)
 - or Toxicological (e.g. AET)

Example: see next slide





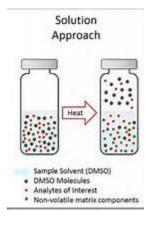
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HS-GC/MS Screening

Volatile Organic Compounds (typically MW < 200)

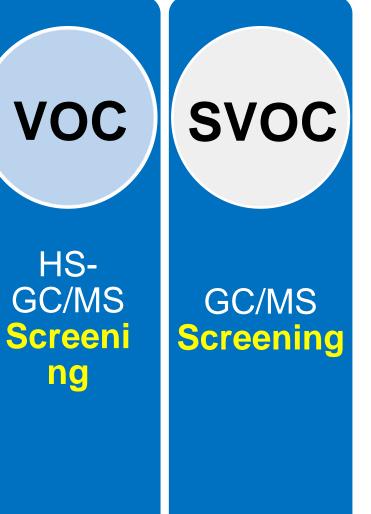
- Monomer Residues
- Solvent Residues from Production steps
- Residues from polymer treatments (e.g. Washing)
- Small Polymer Breakdown products





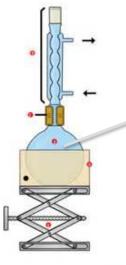


5. ANALYTICAL TECHNIQUES TO PERFORM E/L STUDIES



Semi-Volatile Organic Compounds (MW < 650)

- Lubricants
- Plasticizers
- Antioxidants
- Polymer degradation products
- Solvents with an elevated boiling point







Derivatisation GC/MS

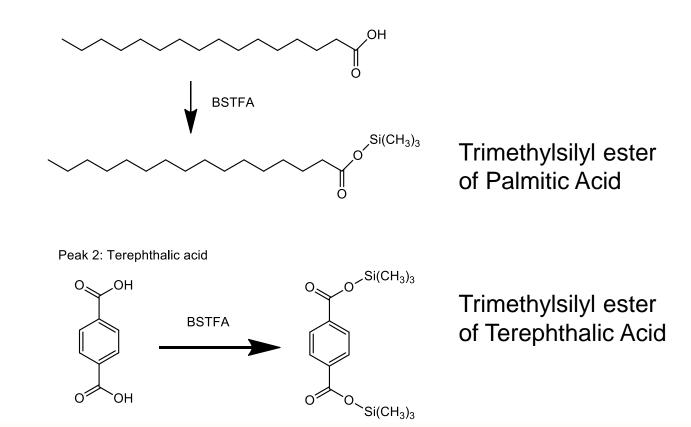
- A combined Headspace-GC/MS, GC/MS and LC/MS approach is suited for a broad list of organic compounds.
- However, compounds containing functional groups such as: Organic acids, Amines, alcohols, polyols, aldehydes, ketones... may not always be very sensitive in regular GC/MS analysis!!
- A Derivatisation Method is using BSTFA as derivatisation agent (conversion to more volatile, less polar trimethylsilyl esters).



ANALYTICAL TECHNIQUES – D.I.-GC/MS

DERIVATISATION GC/MS: EXAMPLES

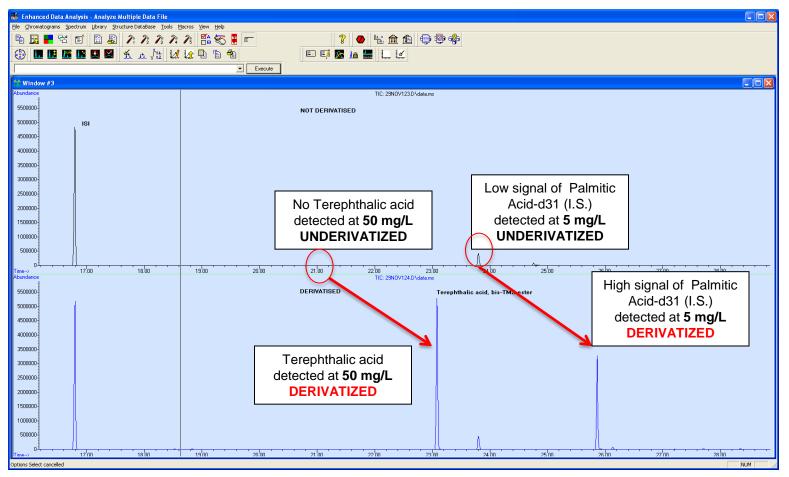
Peak 1: Palmitic acid





ANALYTICAL TECHNIQUES – D.I.-GC/MS

DERIVATISATION GC/MS: RESULTS





ANALYTICAL TECHNIQUES – D.I.-GC/MS

Other GC/MS Techniques (High-End GC/MS)

GC-MS (C.I.): Chemical Ionisation GC/MS

- "Soft Ionization" Compared to Electron Impact (E.I. 70eV)
- The molecule is less Fragmented
- Detection of Molecular Ion
- Allows to determine the Molecular Mass (i.e. With GC-ToF)
- Can be used for "Second Pass" Identifications

GC-QQQ or GC-"Triple Quad" Mass Spectrometer

- **Targeted** analysis in complex matrices
- Very low Detection Limits in complex matrices due to elimination of matrix interferences



Other GC/MS Techniques

GC-(Q)-ToF or GC-"Time-of-Flight" Mass Spectrometer

- Accurate Mass Measurements: what does it bring?
- Principle: Every Atom has a specific Atomic Weight
 - C= 12,00000
 - H = 1,00794
 - O = 15,9994
 - N = 14,0067
 - ...
- Look for the best combination of Atoms which will fit the Accurate Mass the best, Measured with GC-ToF.



GC-TOF Accurate Mass Measurements

Example: a Compound - Accurate Molecular Mass of 136.05243 - was detected.

What could be the Elemental Formula? Using a CALCULATOR

Specify the mass				
Accurate mass experimental result: 136.052430				
Results:				
MF	Monoisotopic mass	PPM	mDa	unsaturation
$1 C_8H_8O_2$	136.0524295014	0.004	0	5
$2 C_3H_7FN_3O_2$	136.0522296921	1.472	-0.2	1.5
3 C ₅ H ₁₁ ClNO	136.0529166949	3.577	0.487	0.5
4 CH ₈ N ₆ S	136.0531149801	5.035	0.685	1
5 C ₃ H ₉ ClN ₄	136.0515740244	6.292	-0.856	1

Most Probably, the Elemental Formula of this molecule is $C_8H_8O_2$

Cross Examining results of other Analytical results, revealed that this compound is **4-methylbenzoic acid**

However, this conclusion cannot be drawn, based solely on accurate mass!



Other GC/MS Techniques

GC-ToF or GC-"Time-of-Flight" Mass Spectrometer

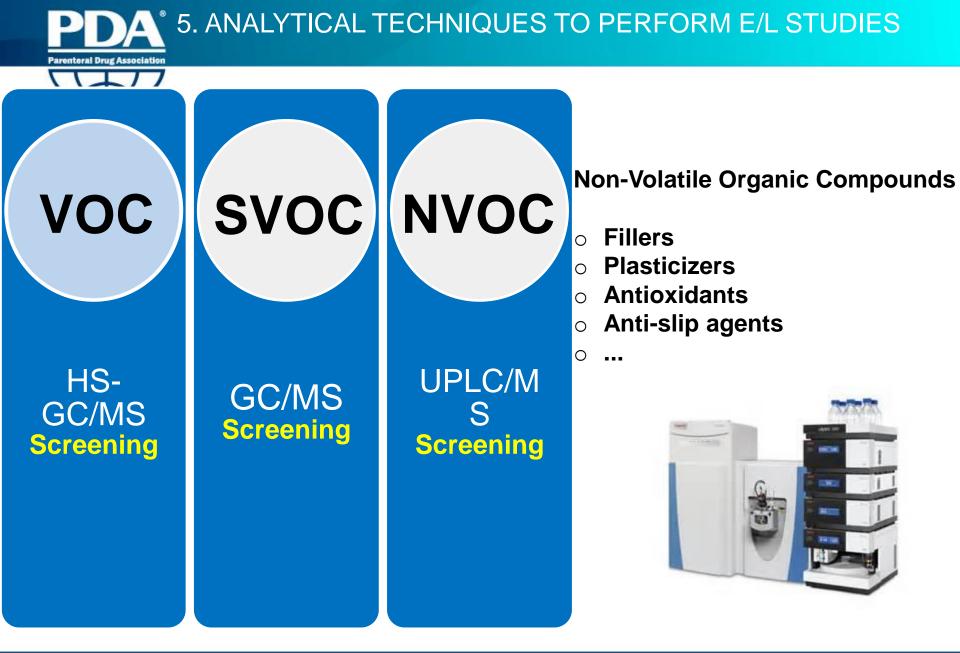
- For extracts with a lot of "Unknown" compounds, the extracts are analyzed with GC-ToF (in E.I. and C.I. Mode) in order to determine the
 - 1. Molecular lon and hence the Elemental Composition (Cl and/or El)
 - 2. Fragment information (EI)
 - 3. In combination with existing data, determine more about the **Structure and Source** of the compound
 - 4. In some cases, in combination with **Derivatization Procedure**
 - 5. In some cases, a full identification of the compound



However: Overlap with compounds from GC/MS (Volatile & Semi-Volatile Compounds)

The principle of HPLC

- High Pressure
- Separation, mostly reverse phase chromatography
- Optimizing separations by
 - Selection of Chromatographic Column (Polarity, Length...)
 - Selection of the Elution Solution (WFI, MeOH, ACN...)
- Detection of the Compounds (UV: DAD; Mass Detection)





* ANALYTICAL TECHNIQUES – LC/MS (UPLC-HRAM)



HPLC - UV

Advantages

- Standard Equipment in a Lab
- o Cost
- UV-Detector can be a *nice addition* to other Detectors, e.g. MS

Disadvantages

- Not a Universal Detector (Target Molecules need Chromophores)
- Non specific
- Not very Sensitive
- $\circ~$ Information about the Detected Molecule is limited
 - $_{\odot}$ E.g. Is the molecule linked to the API?





Advantages

- o Specificity
- o Sensitivity
- More can be said about the Identity of the Compound
- Quality of Information HRAM > Low Resolution
- Allows to build Databases for Identification

Disadvantages

- \circ Cost
- Not a Universal Detector (Target Molecules need to Ionize)
- However, different Ionisation Modes allow a broader detection of Compounds (APCI+/-; ESI+/-)

ANALYTICAL TECHNIQUES – LC/MS (UPLC-HRAM)

LC-MS

Older systems: LOW Resolution Mass spectrometer Ion Trap/Single Quad

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Accuracy of Mass Detection is poor: 1 Dalton

m/z 220 can be distinguished from 221

HIGH Resolution LC-MS (LC-HRAM)

Orbitrap/Time-of-Flight (ToF)

Accuracy of Mass Detection - Orbitrap:

Mass error : sub ppm m/z 220,2456 can be distinguished from m/z 220,2457

MAJOR ADVANTAGES!

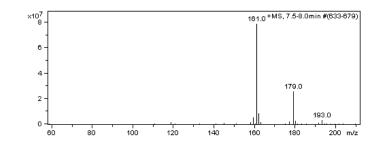
- » Robust: accurate mass is independent of the system
- » High Accuracy in mass detection allows elemental composition analysis of an unknown analyte
- » Extremely powerfull if coupled to a UPLC
- » Building specificity into your databases based on mass accuracy and retention time!

PDA®ANALYTICAL TECHNIQUES – LC/MS (UPLC-HRAM)





LOW RESOLUTION MASS



No information

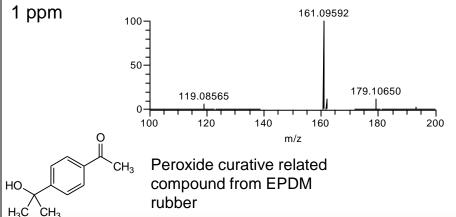
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LC-ORBITRAP (HIGH MASS ACCURACY)



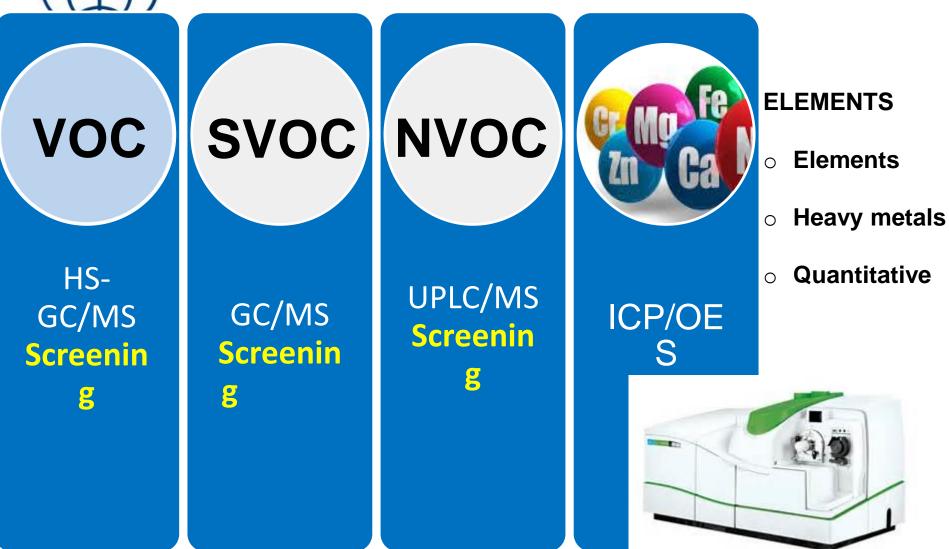
HIGH RESOLUTION ACCURATE MASS

 $C_{11}H_{14}O_2$ exact monoisotopic mass: 179.10666 Mass error:





5. ANALYTICAL TECHNIQUES TO PERFORM E/L STUDIES









ICP-OES or ICP-MS:

- Metals from Glass
- Metals from Rubbers
- Catalysts, used on the polymerization
- Fillers, added to Polymers
- Acid Scavengers

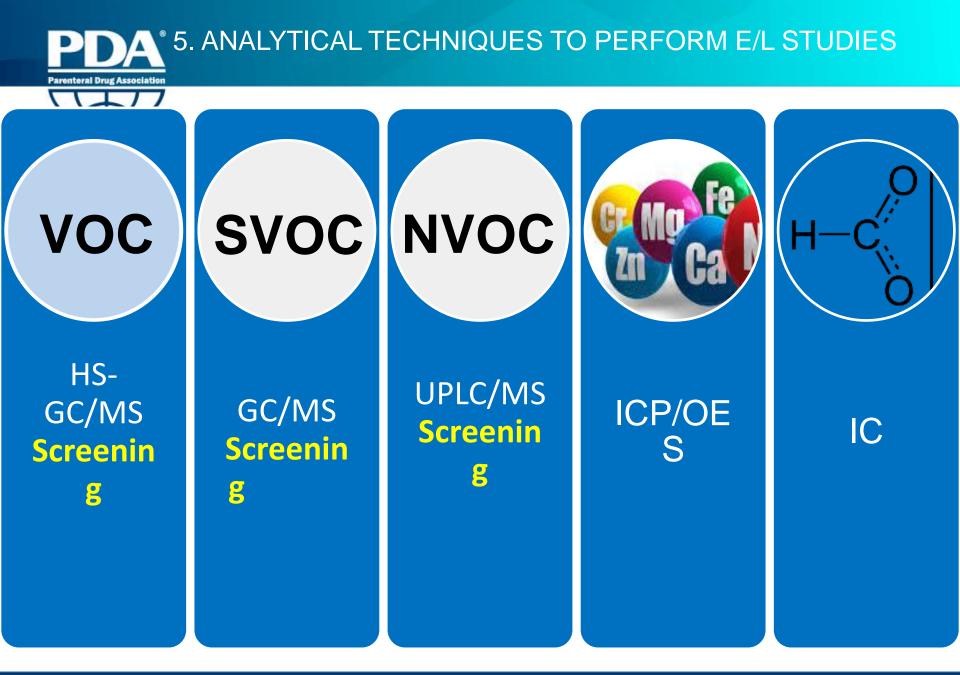
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Activator systems for Rubbers





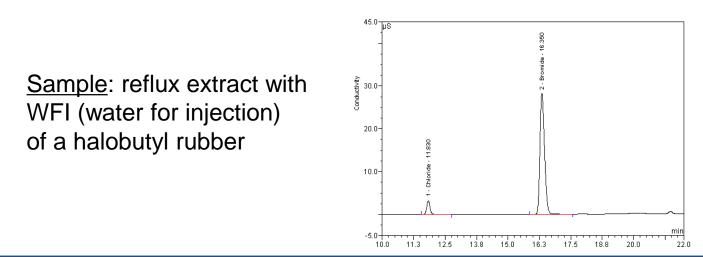
ICP-OES



PDA OTHER TECHNIQUES

Ion Chromatography:

- PolyOlefins (e.g. After Irradiation/Ageing): Acetate & Formate
- Halobutyl Rubbers: Bromide, Chloride, Fluoride
- > Other trace impurities: Nitrite, Nitrate, Phosphate, Sulphate
- <u>Example</u>: Halobutyl rubbers may contain traces of bromide or chloride ions, either from side-products generated during the halogenation step, or rubber degradation products, or impurities. Additionally, fluoride may be released from fluoropolymer coatings





OTHER SPECIFIC METHODS

- ✓ **GF-AAS** For Silicone Oil Detection
- ✓ ESI-UPLC-HRAM (Electron Spray: BPOG Method)
- ✓ HPLC-UV for TMPTMA (glue residue)
- ✓ HPLC-UV for S_8 (Cross Linker)
- ✓ pH (release of acidic/alkalinic agents in UPW)
- ✓ Conductivity (release of salts in UPW)
- ✓ Non-Volatile Residue (gravimetric residue)
- ✓ FTIR characterization of NVR
- Total Organic Carbon: reconsiliation with concentration of organic compounds from chromatographic techniques





ANALYTICAL TECHNIQUES USED FOR LEACHABLES TESTING





TECHNIQUES USED IN LEACHABLE STUDIES

- ✓ Headspace GC/MS: Volatile Compounds
- ✓ Direct Injection GC/MS: Semi-Volatile Compounds
- ✓ D.I. GC-QQQ: Semi-Volatile Compounds
- ✓ LC-QQQ: Non-Volatile Compounds
- ✓ Ion Chormatography: (An)Ions
- ✓ ICP-OES or ICP-MS: Metals

Specific Analysis/Techniques for specific target analyses...

(See further presentation "Leachable Studies")



ANY QUESTIONS?

