

# Water Determination by Karl Fischer Titration



### **Why Water Determination?**

#### Fuel, oils and lubricants

• Water content indicates the danger for corrosion



#### Keeps airplanes safe!

### **Why Water Determination?**

#### **Pharmaceutical products**

• Water content defines the quality and shelf life



#### **Avoids headaches!**

# **Applications for Karl Fischer Titration**

- Chemical industry
- Pharmaceutical industry
- Petrochemical industry
- Power stations
- Plastic industry
- Feed
- Food and beverage
- Paints, Adhesives
- Cosmetic industry
- ... and more

#### → many different samples!



ation
ontrol
vings
afety
alety
elf life

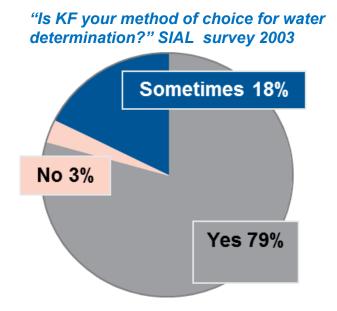
# **Techniques for Moisture/Water Determination**

# Karl Fischer (KF) titration is the most popular water determination technique!

customer survey 2003

#### Alternative techniques

- Weight loss upon drying (most common alternative): Drying cabinet, Microwave drying, Halogen drying, IR drying
- GC/MS
- NIR
- Xylene distillation (DIN ISO 3733)



#### KF titration is always the reference method!

### **Moisture/Water Determination**

- Types of water:
  - water of crystallization (chemically combined)
  - entrapped water
  - adherent moisture

- To determine water of crystallization and entrapped water, sample must be dissolved completely.
- To determine only adherent moisture, dissolution of the sample must be prevented.

### **Why Karl Fischer Titration?**

- Selective: determination of the water content
  - drying techniques cannot differentiate between water and moisture
  - chemically combined water (water of crystallization) may not be detected completely by loss on drying method
- Accurate: very small standard deviations are achievable
- Traceable to water via the standard used for validation
- Wide measuring range: ppm %
- Fast: 1-3 minutes average per determination

# **History of Karl Fischer Titration**

#### The milestones

- 1935 Karl Fischer published a reagent for water determination consisting of iodine, SO<sub>2</sub>, pyridine and methanol
- 1979 Eugen Scholz and Helga Hoffmann, Riedel-de Haën, replaced pyridine by imidazole and invented Hydranal reagents
- 1980 First pyridine-free Hydranal reagents are launched
- 1998 Sigma-Aldrich Laborchemikalien introduced
   the first ethanol-based KF reagents, the Hydranal E-types
- 2008 Hydranal reagents brand change from Riedel-de Haën to Fluka
- 2009 Accreditation according to ISO 17025
- 2014 Accreditation according to ISO 17025 + ISO Guide 34
- 2015 Hydranal became part of Honeywell family









# **Basic Forms of KF Titration**

#### • Volumetric titration:

- with one- or two-component reagents
- water measured in: mg
- water content: 0.01% (100 ppm) 100%

#### Coulometric titration:

- with cell with or without diaphragm
- water measured in: µg
- water content: 0.001% (10 ppm) ca. 5-10%

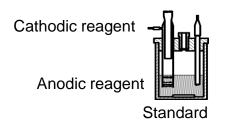
#### • KF titration with an oven:

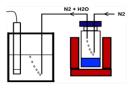
- not a special titration technique
- indirect form of sample introduction
- used in combination with volumetric or coulometric titration



Standard

Titrating agent





### HYDRANAL<sup>™</sup> Karl Fischer Reagents



# **Karl Fischer Titration**

Karl Fischer reaction mechanism:

(1) 
$$CH_3OH + SO_2 + RN \leftrightarrow [RNH]SO_3CH_3$$

(RN = base)

(2)  $H_2O + I_2 + [RNH]SO_3CH_3 + 2 RN \rightarrow [RNH]SO_4CH_3 + 2 [RNH]I$ 

- Reagent for KF titration needs:
  - iodine
  - sulfur dioxide
  - base

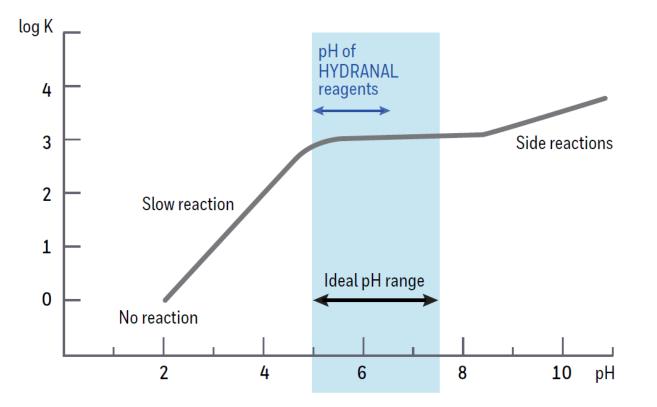
- provide intermediate product
- solvent (alcohol)



- Base  $\rightarrow$  speed, stability, buffering
- Solvent  $\rightarrow$  reactivity, end-point indication, shelf life

#### KF reagent reacts quantitatively and selectively with water

### Influence on pH Value



### **Titer of Reagent**

 $Titer (WE) = \frac{Weight of titrated water [mg]}{Consumption of reagent [mL]}$ 

- Example Titer 5.125: with 1.00 mL consumed reagent, 5.125 mg water have been titrated
- Product Specification:
  - HYDRANAL-Composite 5:
  - HYDRANAL-Titrant 5:
  - HYDRANAL-Composite 2:
  - HYDRANAL-Titrant 2:
  - HYDRANAL-Composite 1:

- titer 4.5-5.5 mg/mL (+/- 10%) → 5.35 mg/mL
- titer 4.95-5.05 mg/mL (+/- 1%)
- titer 1.6-2.4 mg/mL (+/- 20%)
- titer 1.96-2.04 mg/mL (+/- 2%)
- titer 0.8-1.2 mg/mL (+/- 20%)

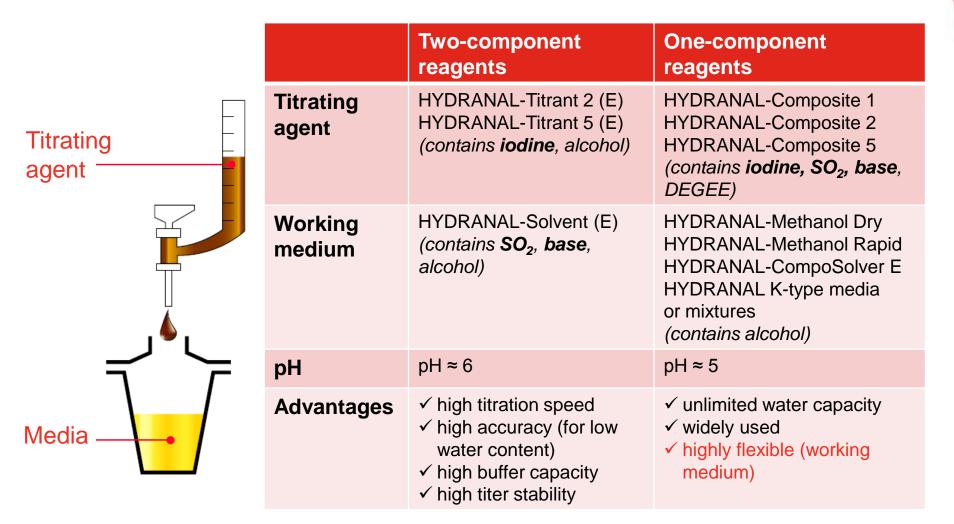
WE = water equivalent

# **Titer of Reagent**

- Influences on titer stability:
  - air humidity
  - temperature change (+1°C  $\rightarrow$  titer change of -0.1%):

example:	Titer 5.050 mg/mL
≻ + 1°C (-0.1%)	Titer 5.045 mg/mL
≻ + 20°C (-2%)	Titer 4.950 mg/mL

# **Volumetric KF Titration**



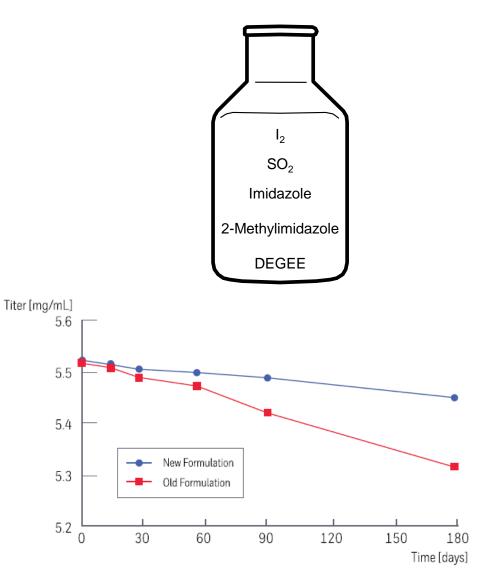
DEGEE = Diethylene glycol monoethyl ether

Highest flexibility by using HYDRANAL-Composite one-component reagents

# **HYDRANAL-Composite**

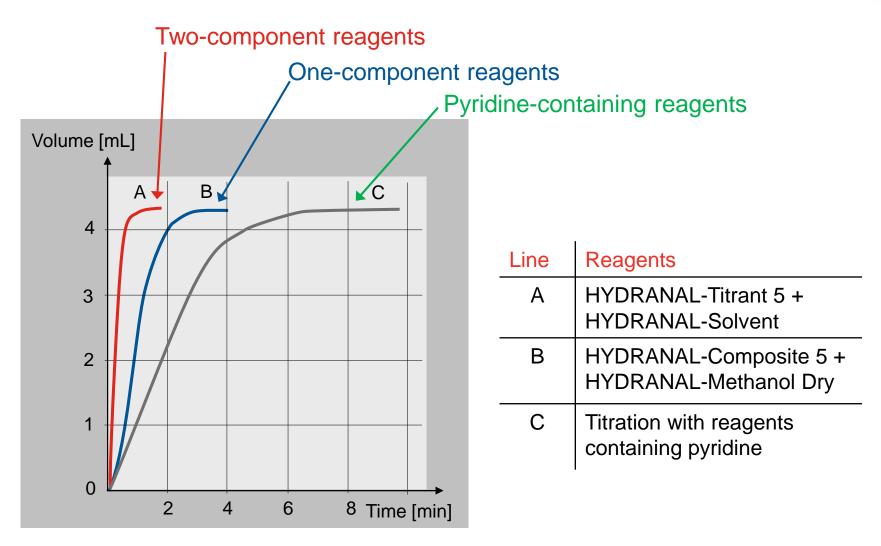
#### Improved quality:

- No crystal formation
- Better stability
- Same performance
- Formulation proprietary
- No methanol



#### DEGEE = Diethylene glycol monoethyl ether

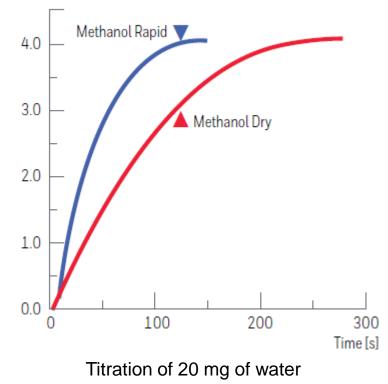
# **Speed of Volumetric KF Titration**



#### Titration of 40 mg of water

### **HYDRANAL-Methanol Rapid**





Advantages:

- Shorten titration time
- Higher accuracy
- Rapid end point

- HYDRANAL-Methanol Dry is not buffered (only methanol)
- HYDRANAL-Methanol Rapid contains accelerators (SO<sub>2</sub>, imidazole)

# **Coulometric KF Titration**

Karl Fischer reaction mechanism:

- (1)  $CH_3OH + SO_2 + RN \leftrightarrow [RNH]SO_3CH_3$  (RN = base)
- (2)  $H_2O + I_2 + [RNH]SO_3CH_3 + 2 RN \rightarrow [RNH]SO_4CH_3 + 2 [RNH]I$
- Iodine electrochemically generated from the oxidation of iodide contained in the coulometric KF reagent

Iodine formation at the anode: $2I^- → I_2 + 2e^-$  (oxidation)Hydrogen formation at the cathode: $2H^+ + 2e^- → H_2$  (reduction)

- Calculation based on the amount of produced (consumed) current over time (acc. to Faraday's law)
- Advantages of coulometric KF titration:
  - easy to use
  - for small amounts of water
  - high accuracy
  - completely glass cell

### Which Cell to Choose?

#### Cell <u>without</u> diaphragm:

- more convenient for the user
- only one reagent required
- less cleaning

#### Cell <u>with</u> diaphragm:

- if reagents contain solubilizer, like: chloroform, xylene, toluene, long chained alcohols
- reagent for ketones (free from methanol)
- in general, more accurate results for small amounts of water (low ppm-range)

# **Coulometric KF Titration**

HYDRANAL- Coulomat (Catholyte)		Reagents preferred for cells with diaphragm	Reagents preferred for cells without diaphragm
	Anolyte	HYDRANAL-Coulomat A* HYDRANAL-Coulomat Oil* HYDRANAL-Coulomat E HYDRANAL-Coulomat AG HYDRANAL-Coulomat AG-Oven HYDRANAL-Coulomat AG-H* HYDRANAL-Coulomat AK*	HYDRANAL-Coulomat AD HYDRANAL-Coulomat E HYDRANAL-Coulomat AG HYDRANAL-Coulomat AG-Oven
HYDRANAL- Coulomat (Anolyte)	Catholyte	HYDRANAL-Coulomat CG HYDRANAL-Coulomat CG-K	Not used

\* higher recovery when used with cell without diaphragm

### **HYDRANAL** Reagents for Oils

Solvents for volumetric titration with one-component reagents

HYDRANAL-Solver (Crude) Oil (contains chloroform, xylene and methanol; fulfills the requirement of ASTM D 4377-00 for water determination in oils and solvents) HYDRANAL-LipoSolver CM (contains chloroform and methanol) HYDRANAL-LipoSolver MH (contains methanol and hexanol)

Solvents for volumetric titration with two-component reagents

HYDRANAL-Solvent CM (contains chloroform and methanol) HYDRANAL-Solvent Oil (contains methanol and hexanol)

#### Anolyte for coulometric titration with diaphragm

HYDRANAL-Coulomat Oil (contains chloroform, xylene & methanol)

Water standard

HYDRANAL-Water Standard Oil (based on mineral oil)

### **HYDRANAL-Water Standards**

- For volumetric and coulometric KF titrations
- Manufactured according to current ISO standards
- Verified against NIST SRM 2890
- Long shelf life (up to 5 years)
- Convenient packaging
- Supplied with detailed instruction and Report of Analysis with exact water content
- Liquid standards are packaged in glass ampoules under argon. Each box contains ten single-use ampoules, easy to open (pre-notched).
- Solid standards are packed in amber glass bottles. They contain defined amounts of chemically combined water.



### **HYDRANAL-Water Standards**

Туре	Name	Form	Water content *
Analytical standards (tested against NIST SRM 2890)	HYDRANAL-Water Standard 10.0 HYDRANAL-Water Standard 1.0 HYDRANAL-Water Standard 0.1 HYDRANAL-Water Standard 0.1 PC <b>NEW!</b> HYDRANAL-Water Standard Oil HYDRANAL-Standard Sodium Tartrate Dihydrate (34696) HYDRANAL-Water Standard KF Oven 140-160°C HYDRANAL-Water Standard KF Oven 220-230°C	Liquid Liquid Liquid Liquid Solid Solid	10.0 mg/g = 1.0% 1.0 mg/g = 0.1% 0.1 mg/g = 0.01% 0.1 mg/g = 0.01% <50 ppm = 0.005% ~15.66% ~5.0% ~5.55%

\* exact value on RoA

Report of	Analysis	Honeyw	ell   Fluka <sup>.</sup>
Analysed for Honeywell Laborchemikalien G	mbH, Supply chain, Wunstorfe	r Str.40, D-30926 Seelze	
Product: HYDRANAL®-Water Standa	rd KF-Oven 140°C-160°C	Cat. No.: 34693	Lot: SZBG076AV
The water content of this lot is:	5.08 % ± 0.02 %	(k=2; 95% confidence	interval)
5 1 2	ter and tested against NIST SR	M 2050.	
Thomas Wendt, Supervisor	Received against wish on	Thomas Wardt	Seelze
Honeywell Laborchemikalien GmbH Thomas Wendt, Supervisor Technical Service HYDRANAL® Wunstorfer Str.40, D-30926 Seelze	Received against wish six		QC release date Seelze 25. May 2016 Page 1 of 1

### HYDRANAL-Water Standard 0.1 PC NEW!

- Based on propylene carbonate
- Improved stability compared to HYDRANAL-Water Standard 0.1:
  - shelf life increased from 2 years to 3 years
  - can be stored at room temp. instead of 2-8°C

### HYDRANAL-CRM Water Standards NEW!

- In 2014, Hydranal Technical Service in Seelze completed its combined accreditation according to ISO/IEC 17025 and ISO Guide 34, the socalled "Gold Standard Accreditation", which is the highest achievable quality level for producers of Certified Reference Materials (CRMs).
- With the double accreditation, Hydranal introduced the very first commercially available CRM Water Standards for Karl Fischer titration.

Туре	Name	Form	Water content *
Certified	HYDRANAL-CRM Water Standard 10.0	Liquid	10.0 mg/g = 1.0%
Reference	HYDRANAL-CRM Water Standard 1.0	Liquid	1.0 mg/g = 0.1%
Materials	HYDRANAL-CRM Sodium Tartrate Dihydrate	Solid	~15.66%

\* exact value on CoA

### **Sample Handling**



### Sample size calculations

Sample size  $[g] = 0.5 \times \frac{Burette's \ volume \ [mL] \times Titer \ [mg/mL]}{Expected \ water \ content \ [mg/g]}$ 

Sample	Sample size	Titrant volume used
100% water = 1000 mg/g	Titer 5: 25-50 mg Titer 2: 10-20 mg	5-10 mL 5-10 mL
15.6% water = 156 mg/g (34696 HYDRANAL-Sodium Tartrate Dihydrate)	Titer 5: 150-200 mg Titer 2: ca. 80 mg	4.7-6.2 mL 6.2 mL
1.0% water = 10 mg/g (34849 HYDRANAL-Water Standard 10.0)	Titer 5: ca. 2 g Titer 2: 1-2 g	4 mL 5-10 mL

Titrant volume used should not exceed burette volume! Ideally: half of burette volume.

### **Sample Size Recommendations**

Titer 5 mg/mL				Titer 2 mg/m	L		Titer 1 mg/m	L		
		E	Burette volum	ne	E	Burette volum	ne	E	Burette volum	ne
		5 mL	10 mL	20 mL	5 mL	10 mL	20 mL	5 mL	10 mL	20 mL
		Recomn	nended samp	le size (g)	Recomn	nended samp	le size (g)	Recomn	nended samp	le size (g)
	90 %	Х	0.04	0.08	0.007	0.015	Х	Х	Х	Х
	75 %	Х	0.05	0.1	0.01	0.02	Х	Х	Х	0.02
ent	50 %	Х	0.08	0.16	0.015	0.03	0.05	Х	0.015	0.025
content	20 %	0.08	0.125	0.25	0.025	0.05	0.1	Х	0.025	0.05
	10 %	0.125	0.25	0.5	0.05	0.1	0.2	0.025	0.05	0.1
water	5 %	0.25	0.5	1	0.1	0.2	0.4	0.05	0.1	0.2
	2.5 %	0.5	1	2	0.2	0.4	0.8	0.1	0.2	0.4
Expected	0.25 %	5	10	20	2	4	8	1	2	4
Exp	0.1 % (1000 ppm)	12.5	25	25	5	10	20	3	6	12
	0.01 % (100 ppm)	25	25	Х	25	25	Х	25	25	Х
	0.001 % (10 ppm)	Х	Х	Х	25	Х	Х	25	Х	Х

Consumption >1/2 burette volume

Consumption approx. 1/2 burette volume

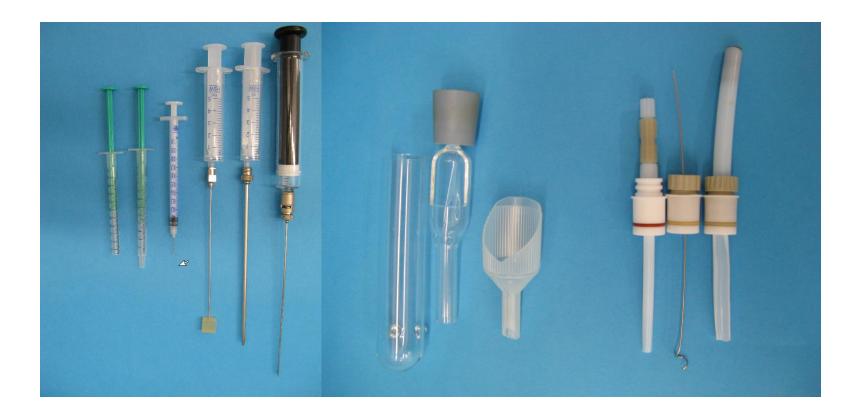
Consumption < 1/2 burette volume

X = not recommended

#### Honeywell

Honeywell Confidential - © 2016 by Honeywell International Inc. All rights reserved.

### **Liquid and Solid Samples Handling**



#### Every sample should be weighed by difference!

Honeywell

### **Liquid Water Standards Handling**

- Standards in 8 mL ampoules → use 10 mL glass syringe
  - HYDRANAL-CRM Water Standard 10.0
  - HYDRANAL-Water Standard 10.0
  - HYDRANAL-Water Standard Oil
- Standards in 4 mL ampoules → use 5 mL glass syringe
  - HYDRANAL-CRM Water Standard 1.0
  - HYDRANAL-Water Standard 1.0
  - HYDRANAL-Water Standard 0.1
  - HYDRANAL-Water Standard 0.1 PC
- Rinse the syringe with 1 mL of water standard and draw the rest of the standard into the syringe in order to protect it against air influences.
- The amount of one ampoule is dedicated for rinsing the syringe followed by a triple determination of the water content.

### **Sample Dissolution**



# **Solutions for Solubility Problems**

Matrix	Variation of titration
Fats, oils, long-chained hydrocarbons	Addition of long-chained alcohols, chloroform or xylene
Proteins, carbohydrates	Addition of formamide
Other insoluble substances	<ul> <li>Titration at 50°C or in boiling methanol</li> <li>Homogenizing device</li> <li>Extraction of water with suitable solvent</li> <li>KF oven method</li> </ul>

## **Sample Dissolution in Volumetric Reagents**

Solubilizer	Sample	Amount	Ratio to methanol
<b>1-Hexanol</b> (long-chained alcohols)	for long chained hydrocarbons, dispersion of oils	max. 50%	(1:1)
Chloroform	for oils, organic samples	max. 70%	(2:1)
Xylene, Toluene	for crude oil, organic samples	max. 70%	(2:1)
<b>Formamide</b> (fresh solution, max. 2 days)	for solids, sugars, proteins	max. 50%	(1:1)

DMSO (dimethyl sulfoxide) – not recommended, alters the KF reaction, results are too low

#### Honeywell Confidential - © 2016 by Honeywell International Inc. All rights reserved.

### **HYDRANAL Auxiliary Reagents**

#### **For dissolution**

HYDRANAL-Formamide dry (max. 0.02% H<sub>2</sub>O) HYDRANAL-Chloroform (max. 0.01% H<sub>2</sub>O) HYDRANAL-Xylene (max. 0.01% H<sub>2</sub>O)

# **Sample Dissolution in Coulometric Reagents**

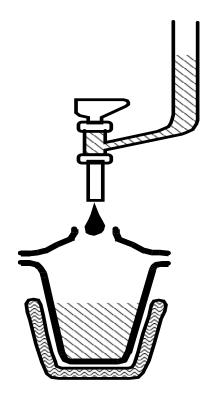
- Coulometric reagents with solubilizing agents (based on methanol):
  - HYDRANAL-Coulomat A (contains chloroform)
  - HYDRANAL-Coulomat Oil (contains xylene and chloroform)
  - HYDRANAL-Coulomat AG-H (contains long-chained alcohols)
- Formamide only limited
  - cell without diaphragm
  - external sample treatment
  - no mixture of reagent and formamide

### **Titration at Elevated Temperature**

 Titration at 40-50°C: double-walled titration cell (connected to water bath)

• Titration in boiling methanol:

for volumetric one-component reagents with reflux condenser and heating mantle (custom-made titration cell required)



## **Homogenizing Device**

- High speed stirrer
- Homogenizes the sample
- Better dissolution and extraction
- Shortens analysis time
- Use in addition to solubilizers and heat







#### **Water Extraction**



### **Internal** Water Extraction (Sample Dissolution)

Solubilizer	Sample
<b>1-Hexanol</b> (long-chained alcohols)	for long chained hydrocarbons, dispersion of oils
Chloroform	for oils, organic samples
Xylene, Toluene	for crude oil, organic samples
Formamide	for solids, sugars, proteins

#### **External** Water Extraction

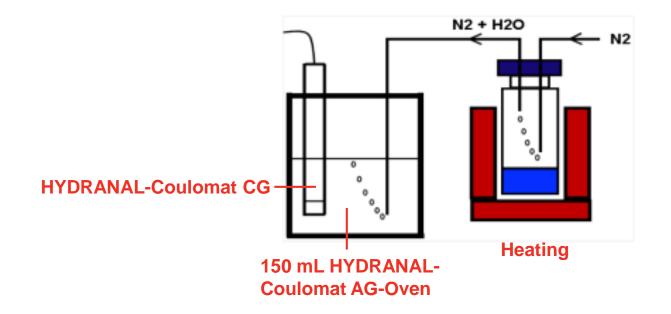
- Samples releasing water very slowly
- Very inhomogeneous samples large sample masses required
- Procedure:
  - general rule: 1 mL of dry methanol will extract approx. 1 mg of water
  - calculate volume of dry solvent needed
  - weigh the sample
  - weigh the solvent added
  - prepare **blank** as above
  - stir for few hours
  - take an aliquot and titrate



### Karl Fischer Oven for Coulometric and Volumetric Titration



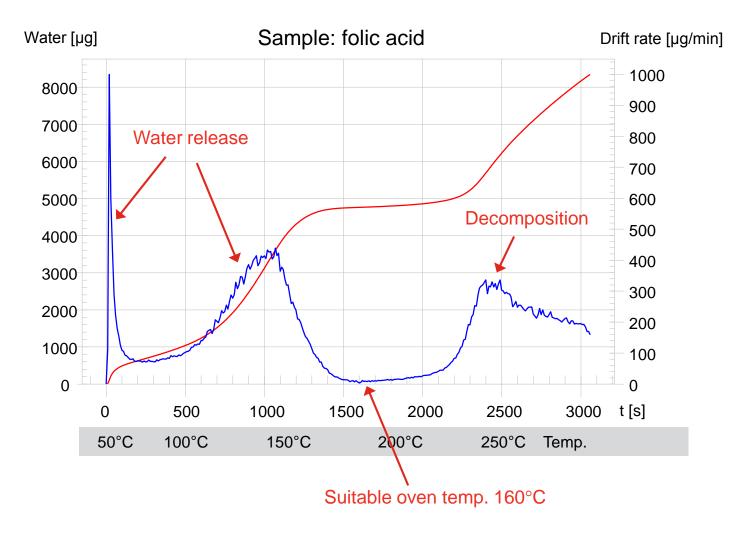
#### **Karl Fischer Oven**



### Karl Fischer Oven

- Ideal reagent is 34739 HYDRANAL-Coulomat AG-Oven with especially low drift:
  - methanol evaporates from the titration cell through the carrier gas
  - in Coulomat AG-Oven methanol is partially substituted with propylene glycol for better stability of drift and overall performance
  - at the end of a working day the original level shall be refilled by addition of dry methanol
- Carrier gas (dried by passing through molecular sieves):
  - air
  - nitrogen, argon for samples sensitive to oxygen
- Ideal heating temperature for the sample:
  - sample water should be released as fast as possible, but decomposition of the sample must be avoided

#### **Temperature Ramp for KF Oven**



44

#### Honeywell Confidential - © 2016 by Honeywell International Inc. All rights reserved.

### **KF Oven Temperature**

Sample	Oven temperature
Biodiesel	100°C
Oils containing additives	80-140°C
Folic acid	160°C
Polypropylene	160-180°C
Polycarbonate	180°C
Polyamide 66	200°C
Zinc peroxide	200°C
Calciumsulfate dihydrate	300°C
tri-Magnesium phosphate	300°C

#### Troubleshooting: Side Reactions



### **Side Reactions with Ketones and Aldehydes**

 Aldehydes and ketones react with methanol to form acetals and ketals respectively – reaction forms water, which is also titrated, resulting in vanishing end points and incorrectly high water content

$$\begin{array}{c} R \\ C = 0 + \begin{array}{c} CH_{3}OH \\ CH_{3}OH \end{array} \end{array} \xrightarrow{R} \begin{array}{c} OCH_{3} \\ C \\ C \\ OCH_{3} \end{array} \xrightarrow{H_{2}O} \end{array}$$

 Additionally, aldehydes can cause a second side reaction, the bisulfite addition, which consumes water and leads to incorrectly low water content

$$\begin{array}{c} H \\ C = 0 + SO_2 + H_2O + NR' \end{array} \xrightarrow{H} \begin{array}{c} SO_3 HNR' \\ R \end{array}$$

### **HYDRANAL-K Type Volumetric Reagents**

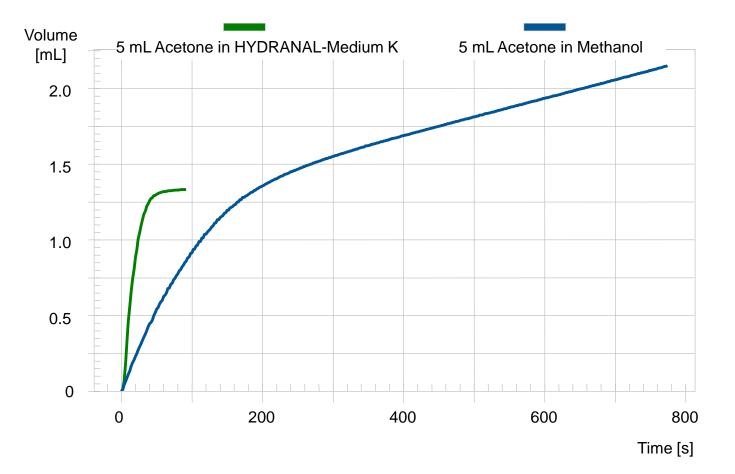
 Specially designed for water determination in aldehydes and ketones without side reactions

Volumetric titrating agents	Volumetric working media
HYDRANAL-Composite 5 HYDRANAL-Composite 5 K	HYDRANAL-Medium K HYDRANAL-KetoSolver HYDRANAL-Working Medium K

- HYDRANAL-Composite 5 can be used with ketones (no methanol)
- HYDRANAL-Composite 5 K is less reactive than Composite 5, can prevent bisulfite addition to a large extend, so should be used with aldehydes (also useful for some ketones)
- HYDRANAL-Medium K is less toxic working medium should be the first choice.

### **Influence of Working Media for Ketones**

Titration of 5 mL acetone with HYDRANAL-Composite 5 in different working media



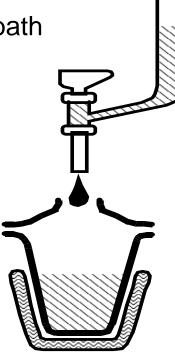
### **HYDRANAL-K Type Coulometric Reagents**

- Coulometric titration should be used for ketone samples only, it is not advised for reactive aldehydes
- Only cells with diaphragm should be used

	Coulometric reagent – catholyte
HYDRANAL-Coulomat AK	HYDRANAL-Coulomat CG-K

### **Titration at Low Temperature**

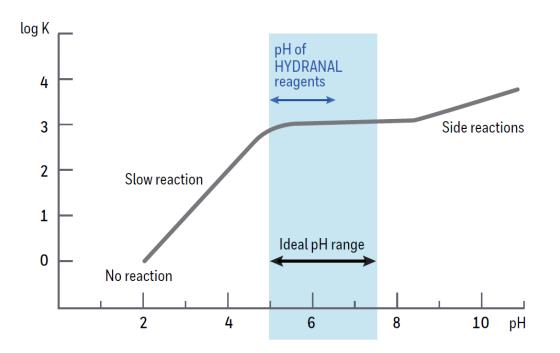
- Method to supress slow side reactions
- Titration vessel placed in an ice bath or double-walled titration cell connected to cooling bath
- Recommended only for volumetric titration:
  - HYDRANAL-Solvent/Titrant: max. -60°C
  - HYDRANAL-Composite: max. -20°C (due to increasing viscosity)



#### Troubleshooting: Neutralization



### **Neutralization of Acids and Bases**



- Checking the pH value:
  - alcoholic medium  $\rightarrow$  only rough check of pH value
  - use pH electrode or pH paper (add deionised water)

#### **HYDRANAL Auxiliary Reagents**

#### **For neutralization**

HYDRANAL-Buffer Acid (buffer capacity 5 mmol acid / mL) HYDRANAL-Buffer Base (buffer capacity 1 mmol base / mL) HYDRANAL-Imidazole (max. 0.1% H<sub>2</sub>O) HYDRANAL-Benzoic acid (max. 0.2% H<sub>2</sub>O) HYDRANAL-Salicylic acid (max. 0.2% H<sub>2</sub>O)

#### **Neutralization of Acids and Bases:** Modifications of Working Media

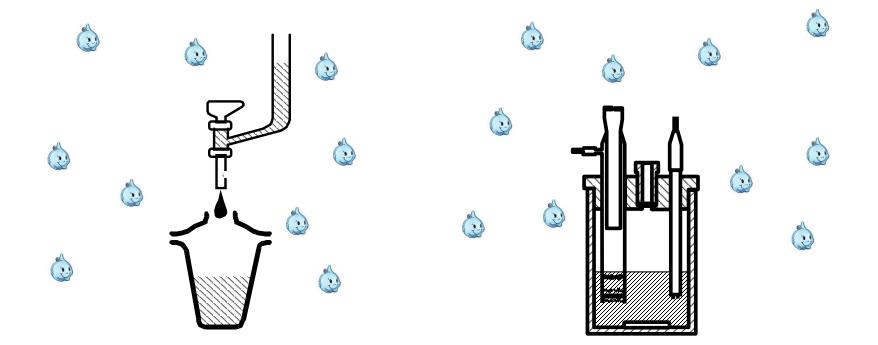
One-component volumetric titration	Two-component volumetric titration	Coulometric titration (cell with diaphragm)
Neutralization of acids		
Add (1:2) HYDRANAL-Buffer Acid or use alone	Add HYDRANAL-Imidazole (7 g/30 mL) or use HYDRANAL-Buffer Acid	Add HYDRANAL-Imidazole (20 g/100 mL)
Neutralization of bases		
Add HYDRANAL-Benzoic Acid or HYDRANAL-Salicylic Acid (5-7 g/30 mL) or use HYDRANAL-Buffer Base	Add HYDRANAL-Benzoic Acid or HYDRANAL-Salicylic Acid (5-7 g/30 mL)	Add HYDRANAL-Benzoic Acid (20 g/100 mL)

**Note:** Use of imidazole with pure methanol in one-component volumetric titration is not recommended (shifts pH too far).

#### Tips & Tricks: Drift Minimizing



### Humidity of Air



At 20°C and a relative air humidity of 60%, 1 L of air contains 12 mg of water

# **Drying Agents for KF Titration**

- Every reagent bottle and every titration vessel needs a drying tube (also important for improving titer stability)
- HYDRANAL-Humidity Absorber
  - beads, with indicator, water absorption capacity < 25%
  - amorphous alumina silica gel
  - color change from orange to almost colorless
  - regeneration at 140°C (until color comes back)
- HYDRANAL-Molecular Sieve 0.3 nm
  - beads, without indicator, water absorption capacity > 15%
  - regeneration at max. 300°C, min. 4 h





### **Drift in the Volumetric Cell**

- Definition of drift: volume of reagent (iodine) permanently consumed
  - humidity from ambient air penetrates into the cell
  - in volume / time [µL/min] or water amount / time [µg/min]
- Recommended acceptable maximum drift (start of titration): 10 µL/min
- End-point criterium: drift < 20 μL/min</li>
- Drift calculations:

$10 \mu L/min = 0.01 mL$ reagent titer 5	$\rightarrow$	50 µg water / min
$10 \mu L/min = 0.01 mL$ reagent titer 2	$\rightarrow$	20 µg water / min
$10 \mu L/min = 0.01 mL$ reagent titer 1	$\rightarrow$	10 µg water / min

- Accepted drift means accepted amount of water migrating into the titration vessel – the cell is stable for using with that titrant
- Cell with reagent titer 1 needs perfect conditions of titration vessel
- New user should always start with reagent titer 5

### **Drift in the Coulometric Cell**

• After fresh refill with HYDRANAL-Coulomat:

- after short time (15 min): < 10 µg/min</p>
- ideally: < 5 µg/min</p>
- often: < 2 µg/min</p>
- After longer use of reagent:
  - max. 20 µg/min (e.g. when determining ketones)
- Unsteady, high drift value  $\rightarrow$  change of reagent necessary

### **Coulometric Cell - Anolyte**

- Anodic compartment is filled with approx. 100 mL anolyte reagent
- Change reagent when:
  - maximum sample volume is reached (cell is full!) = 100 mL reagent plus max. 50 mL sample
  - unsteady or increasing drift
  - water capacity limit is reached: 100 mL anolyte has a capacity of 1'000 mg water (= 1'000'000 µg)
- The same rules for cell without diaphragm

### **Coulometric Cell - Catholyte**

- Moisture in the cathodic compartment will not be eliminated (no KF reaction in the cathodic compartment!)
- Catholyte used for > 1 week:
  - formation of lower sulfur compounds
  - reaction with iodine, diffusion through diaphragm, oxidation at the anode
  - high drift and contaminated diaphragm
- After several weeks:
  - cathode turns black and cathodic compartment starts to smell
  - precipitations / crystal formation may occur

### **Drift with KF Oven**

- Low and stable drift needed
- Drift is subtracted automatically for the duration of determination
- Constant flow rate of the dried carrier gas (air, nitrogen, argon)
  - 120-150 mL/min for tube furnaces
  - 70-80 mL/min for vial oven
- Ideal reagent:
  - HYDRANAL-Coulomat AG-Oven with especially low drift
- Ideal heating temperature:
  - to release the water quickly, but without decomposition of the sample

#### Tips & Tricks: Safer Reagents



### **New Classification According to GHS**

- New and more stringent evaluation of chemicals by the European Chemicals Agency (ECHA) – imidazole classified as a dangerous component
- New guidelines defining how to safely handle HYDRANAL products
- According to the new European Regulation on Classification, Labeling and Packaging of chemical substances and mixtures (CLP):
  - reagents containing imidazole of 0.3% w/w or more must show the following Pictograms and Hazard statements:
    - GHS08 (Health Hazard)
    - H360D (May damage the unborn child)



### **New Classification According to GHS**

- Newly measured values for imidazole concern oral intake of the substance, whereas values for dermal and inhalation remain unchanged
- Karl Fischer reagents are handled in closed systems
  - direct contact with the chemicals is typically prevented and can only occur accidentally
- Hydranal reagents contain imidazole only in dissolved and diluted form
- Imidazole in its free form is only contained to a low degree
  - risk of exposure to inhalation of powder dust is negligible
  - use of personal protection is advised, especially gloves (avoid skin contact)
  - an oral intake is quite unlikely

### **HYDRANAL-E** Type Reagents

- Based on ethanol instead of methanol less dangerous for user and environment
- Improve solubility of long-chained hydrocarbons especially in HYDRANAL-CompoSolver E
- Ketones like acetone can be titrated in HYDRANAL-CompoSolver E (not possible in methanol-based working media)
- Corresponding to regulations of ISO 14001 (environmental management)

# Green HYDRANAL Reagents Based on Ethanol / DEGEE

One-component volumetric reagents	Two-component volumetric reagents	Coulometric reagent
HYDRANAL-Composite 1 HYDRANAL-Composite 2 HYDRANAL-Composite 5 HYDRANAL-Composite 5 K HYDRANAL-CompoSolver E	HYDRANAL-Titrant 2 E HYDRANAL-Titrant 5 E HYDRANAL-Solvent E*	HYDRANAL-Coulomat E

\* should be used quickly after opening

### Applications: Pharmaceutical Industry



### **Pharmacopeia Suitability Test**

#### 2.5.12. Semi-Microdetermination (Volumetric titration)

#### **Method A**

"Introduce into the titration vessel *methanol R*, or the solvent indicated in the monograph or recommended by the supplier of the titrant. Where applicable for the apparatus used, eliminate residual water from the measurement cell or carry out a pre-titration. Introduce the substance to be examined rapidly and carry out the titration, stirring for the necessary extraction time."

#### **Method B**

Back titration (basically same requirements)

#### Suitability

"The accuracy of the determination with the chosen titrant must be verified for each substance to be examined. [...] The water content of the substance to be examined is determined using the reagent/solvent system. Thereafter, sequential known amounts of *water R* are added in an appropriate form (at least 5 additions) and the cumulative water content determined after each addition."

#### **Problems with Ph.Eur. Protocols**

Sample	Ph. Eur.	Troubleshooting
Calcium acetate	+ 2 mL Acetic acid	5 g Salicylic acid
D(-) Fructose	Methanol	+ 10 mL Formamide
D(+) Glucose	Methanol	+ 10 mL Formamide
Gentamicin sulfate	Methanol	+ 10 mL Formamide
Ethyl acetate	Titer 5	Titer 2
DMSO	Recovery rate 75%	no solution

Suitability tests are available on request (hydranal@honeywell.com)

## **Suitability Test Protocol Example**

Suitability test according to Ph.Eur., method 2.5.12 Water semi-micro determination

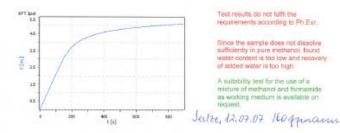
Water determination by	Karl Fischer titration u	ising HYDRANAL®- Composite

Product	Gentamicin sulphate	
Titrant	HYDRANAL®-Composite 5	Titer 5.272 mg/mL
Working medium	30 ml HYDRANAL®-Methanol dry	
Sample handling Procedure	By means of a powder funnel. Weigh by difference The working medium is placed in the titration vest titrant.	
	Then the sample is added and titrated in the same	e way to a stable end point.
Recovery of water added	After achieving the end point, sequential known a titrated in the same way.	mounts of water are added and

	Sample	Water ad	ded			
		1	2	3	4	5
Sample size (g)	0.2857					
Water added (mg)		27.77	28.67	27.85	27.83	27.85
Water found (mg)	26.8466	32.13	28.46	27.85	27.88	27.91
Water content (%)	9.3971					
Recovery (%)		115.71	99.28	100.03	100.16	100.22

#### The reagent/solvent system is considered to be acceptable if:

- The mean recovery is between 97.5% and 102.5%	mean recovery (%)	103.08
- The slope b is between 0.975 and 1.025 (deviation +/- 2.5 %).	slope	1.000
- The error e1 and e2 are not greater than 2.5%	error 1 (%)	15.92
	entor 2 (%)	15.97



#### Suitability test according to Ph.Eur., method 2.5.12 Water semi-micro determination

Water determination by Karl Fischer titration using HYDRANAL®- Composite

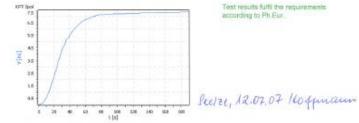
Product	Gentamicin sulphate
Titrant	HYDRANAL®-Composite 5 Titer 5.272 mg/ml.
Working medium	20 ml HYDRANAL®-Methanol dry + 20 ml HYDRANAL®-Formamide dry
Sample handling Procedure	By means of a powder funnel. Weigh by difference. The working medium is placed in the titration vessel and titrated to dryness with the titrant.
	Then the sample is added and titrated in the same way to a stable end point.
Recovery of water added	After achieving the end point, sequential known amounts of water are added and titrated in the same way.

	Sample	Sample Water added					
		1	2	3	4	5	
Sample size (g)	0.3363						
Water added (mg)		38.55	39.00	38.56	39.11	21.16	
Water found (mg)	38.6723	38.36	38.78	38.22	38.62	20.90	
Water content (%)	11.5004						
Recovery (%)		99.49	99.43	99.11	98.74	96.79	

#### The reagent/solvent system is considered to be acceptable if:

- The mean recovery is between 97.5% and 102.5% - The slope b is between 0.975 and 1.025 (deviation +/- 2.5 - The error e1 and e2 are not greater than 2.5%

	mean recovery (%)	99.11	
.5 %).	slope	0.990	
	error 1 (%)	0.69	
	error 2 (%)	1.68	



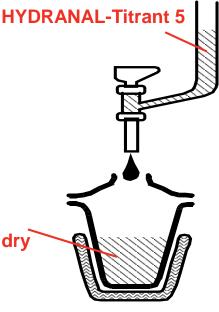
Test results fulfil the requirements

# **Application example: Aspartic acid (L 010)**

- At room temperature, the inherent moisture of the magnesium salt of aspartic acid and the moisture in the magnesium salt of aspartic acid hydrochloride is released so slowly that only a titration at 50°C is possible.
- The magnesium salt of aspartic acid also requires the presence of formamide as solvent agent. Formamide also improves the solubility of the magnesium salt of aspartic acid hydrochloride.

#### Procedure:

- Two-component volumetric titration at 50°C
- Sample size: approx. 0.3 g aspartic acid magnesium salt or 0.15 g aspartic acid magnesium salt hydrochloride



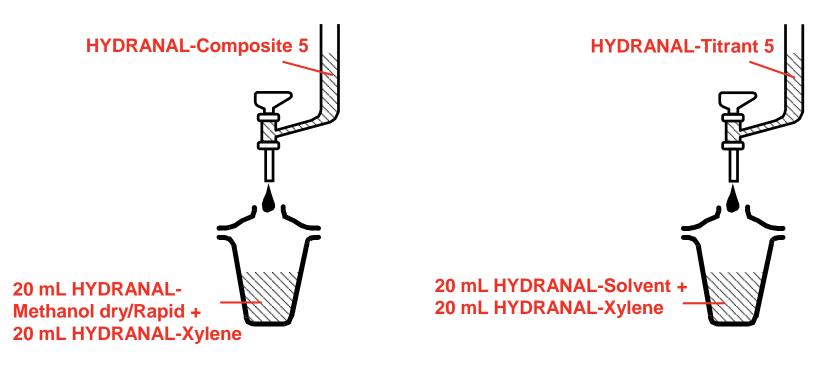
25 mL HYDRANAL-Solvent + 10 mL HYDRANAL-Formamide dry

Heating to 50°C

# Application example: Magrocol 8000 (L 633)

• This sample does not dissolve easily in the alcohol-based KF media. To ensure that it dissolves completely, it is recommended that **xylene** is added.

- One- or two-component volumetric titration
- Sample size: 2 g

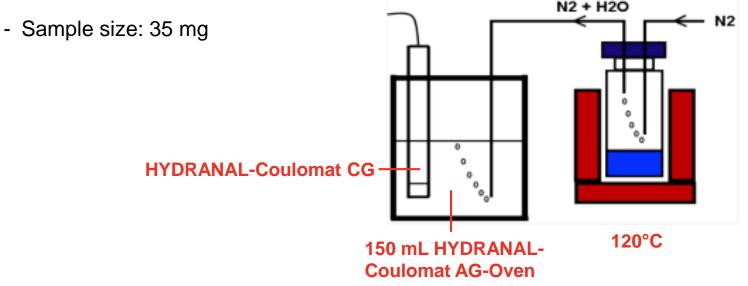


# Application example: Multivitamin (L 404)

- Tablets and powder
- Problems:
  - solubility in methanol (addition of formamide and heating to 50°C didn't give reproducible results)
  - side reaction with iodine

#### Procedure:

Coulometric titration with KF oven



# Application example: Vitamin B12 (L 711)

### • Problems:

- hygroscopicity (weight increase: 1% after 30 s, 7% after 2 h in the air)
- solubility (time to dissolve **150 mg sample**: 3 min./Methanol dry, 1.5 min./Methanol Rapid and 1 min./Solvent)

### Procedure:

	One-component volumetric titration	Two-component volumetric titration	Coulometric titration with KF oven
Titrating agent	Hydranal-Composite 5 or 2	Hydranal-Titrant 5 or 2	-
Working medium	30 mL Hydranal-Methanol Dry or Rapid	30 mL Hydranal-Solvent	150 mL Hydranal-Coulomat AG-Oven (anolyte)
Sample size	0.15 g	0.15 g	0.05 g
Temperature	RT	RT	140°C

# **Application example: Toothpaste (L 029)**

- Toothpaste releases its water content so slowly that a determination is not possible at RT. At 50°C titration still takes ca. 30 min, with added formamide about 15 min.
- With a **homogenizer** the sample is dispersed within few seconds, titration time is only 1.5 min (in methanol).
- Direct titration only if the toothpaste does not contain any carbonates (they might react in the acidic KF medium producing water).

#### Procedure:

- Volumetric titration (high water content: ~40%)
- Sample size: 60-100 mg
- The toothpaste has to be homogenized before the sample is taken
- Sample handling: 1 mL syringe without needle

#### Honeywell Confidential - © 2016 by Honeywell International Inc. All rights reserved.

# **Application example: Toothpaste (L 029)**

	One-component volumetric titration		Two-compone titration	Volumetric titration	
	with homogenizer	without homogenizer	with homogenizer	without homogenizer	with KF oven
Titrating agent	Hydranal- Composite 5	Hydranal- Composite 5	Hydranal- Titrant 5	Hydranal- Titrant 5	Hydranal- Composite 5
Working medium	40 mL Hydranal- Methanol Dry or Rapid	20 mL Hydranal- Methanol Dry or Rapid + 10 mL Hydranal- Formamide dry	40 mL Hydranal- Solvent	20 mL Hydranal- Solvent + 10 mL Hydranal- Formamide dry	40 mL Hydranal- Methanol Dry
Homo- genizer speed	5,000 r/min	-	5,000 r/min	-	-
Tempe- rature	RT	RT	RT	RT	140°C

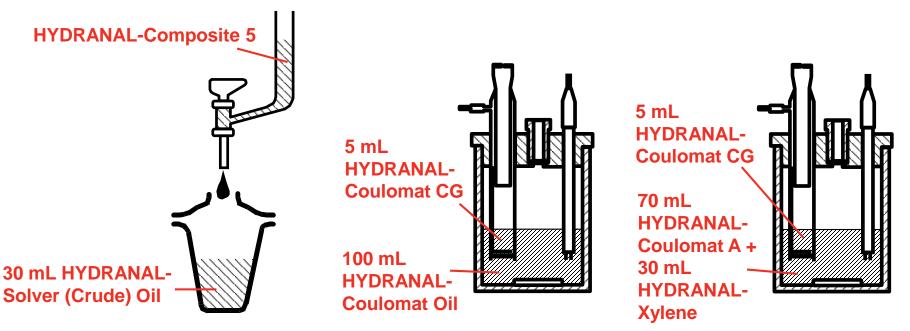
## Applications: Petrochemical Industry



# Application example: Crude Oil (L 108, L 148)

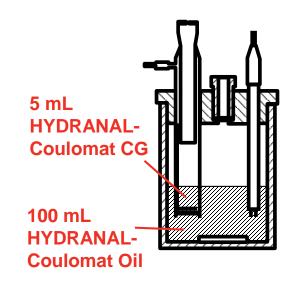
- Intensive homogenisation of samples is a needed for reproducible results.
- Crude oil requires: chloroform to dissolve the oil and xylene to dissolve the tar components. If the tar is not finely dispersed, it can coat the electrode which leads to indication problems.

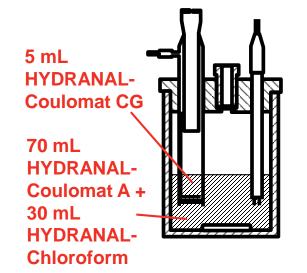
- One-component volumetric or coulometric titration
- Sample size: 4 g for volumetry, 1-2 g for coulometry



# **Application example: Kerosene (L 112)**

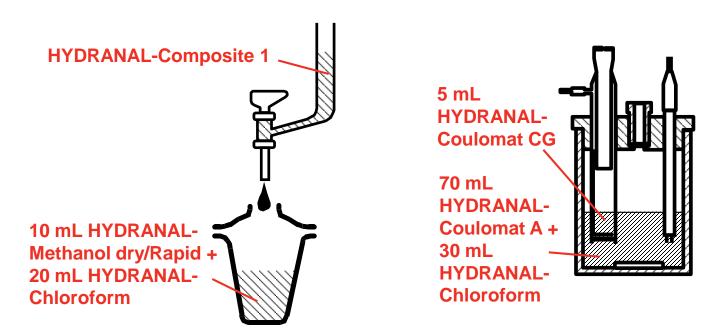
- Low water content (few ppm) only coulometric determination.
- Addition of **chloroform** is needed to improve the solubility of kerosene.
- Procedure:
  - Coulometric titration
  - Sample size: 1-5 mL





# **Application example: Silicon Oil (L 113)**

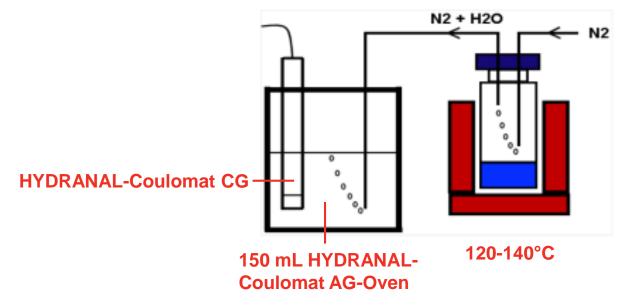
- An additional solvent is necessary to dissolve the sample chloroform.
- Due to the low water content of the material, large samples must be used for volumetric determination.
- Procedure:
  - One-component volumetric or coulometric titration
  - Sample size: 10 mL for volumetry, 1 mL for coulometry



# **Application example: Engine Oil (L 201)**

 The determination of water in engine oil is marked by the presence of very noticeable side reaction. Not only does a side reaction take place with methanol, but also with iodine. The side reaction occurs most strongly in a methanolic working medium. If a non-methanolic working medium is used, the reagent consumption is drastically reduced, yet a significant side reaction is still noticeable by the sluggish titration.

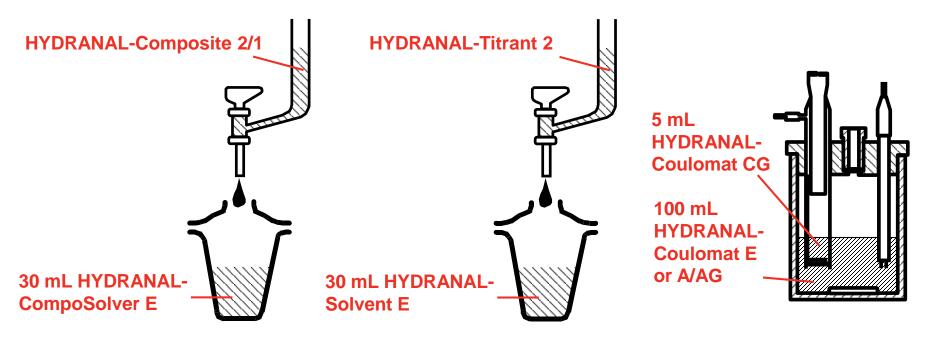
- Coulometric titration with KF oven
- Sample size: 10 g (practical maximum: 2 g)



# Application example: Petrol, unleaded (L 428)

 The solubility of petrol in the methanolic medium of the Karl Fischer titration is limited. Ethanol-based reagents are preferred for volumetric titrations because petrol dissolves well in them. 20 mL petrol dissolve in 30 mL Hydranal-CompoSolver E or Hydranal-Solvent E and 35 mL dissolve in 100 mL Hydranal-Coulomat E.

- Volumetric or coulometric titration
- Sample size: 10 mL for volumetry, 1-2 mL for coulometry



## Applications: Chemical Industry

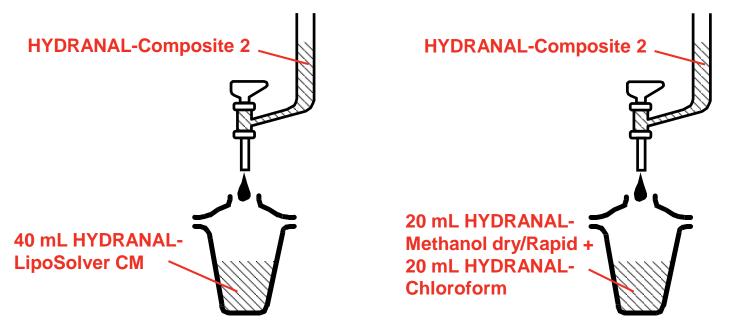


## **Application example: Cyanoacrylate Adhesive (L 118)**

- Cyanoacrylate based adhesives form **lumps** in the alcohol solvents of the KF reagents. The alcohol must be complemented by a solubilizer, e.g. **chloroform**.
- The water content of those adhesives is very low. It is therefore important that the titration cell is adequately very dry before adding the first sample to the medium in the titration vessel.

#### Procedure:

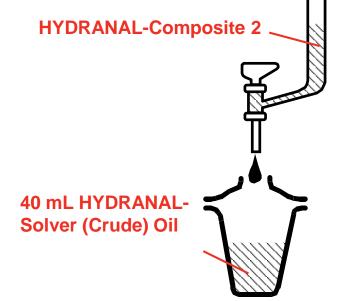
- One-component volumetric titration
- Sample size: 0.5 g

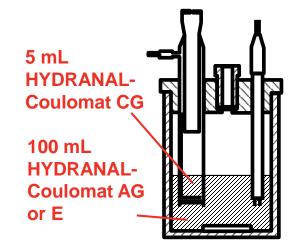


## Application example: Expandable Polystyrene (L 691)

• Expandable polystyrene cannot be determined directly due to **poor solubility** in conventional KF media. This sample must be **dissolved externally** in a suitable solvent, then an aliquot of the solution can be determined in the Karl Fischer cell.

- One-component volumetric or coulometric titration
- Sample preparation: 3 g sample in 50 mL chloroform, stirred 3 h
- Sample size: 5 mL aliquot
- KF oven with coulometer: 220°C in nitrogen. Adapt the size of the sample vessel for the anticipated sample expansion!



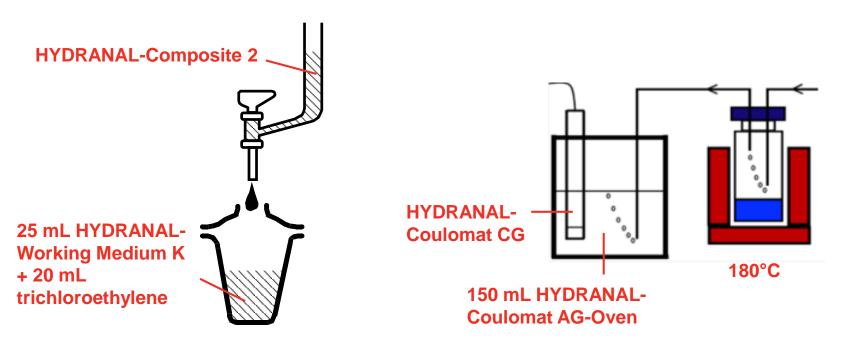


## **Application example: Polycarbonate (L 127, L 129)**

 Due to its poor solubility, the indirect process with a KF oven should be preferred. Polycarbonate is only soluble in certain solvents and precipitates out as flakes from methanolic KF media on the electrodes. It is therefore necessary to carry out the titration using a modified solvent, but it takes 15 minutes for the sample to dissolve.

### Procedure:

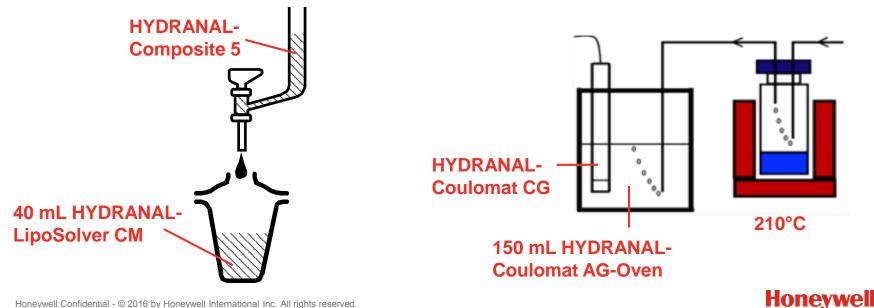
- One-component volumetric or coulometric titration with KF oven
- Sample size: 2 g



## Application example: Poly-L-lactate, PLA (L 577)

 The granulate cannot be dissolved for direct titration in the alcohol-containing media of the KF reagents. This is true even when the media contain portions of chloroform.

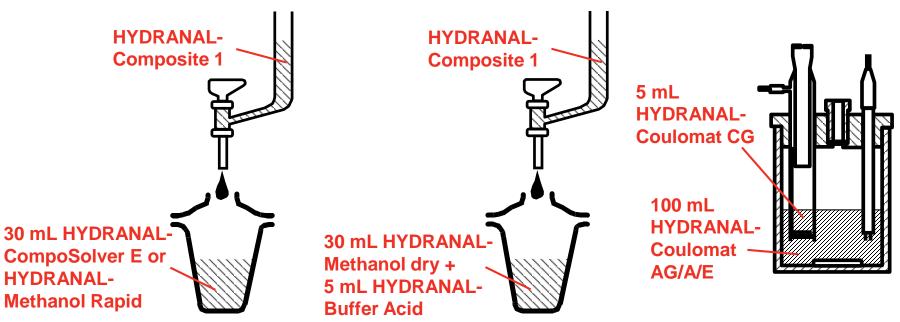
- One-component volumetric or coulometric titration with KF oven
- Sample preparation: 9 g sample in 50 mL chloroform, stir until dissolved to form a viscous solution
- Sample size: 5 mL aliquot
- Sample handling: disposable syringe without a needle
- KF oven with coulometer: 210°C, 0.5 g sample



# **Application example: Caprolactone (L 410)**

 A side-reaction is taking place. If the working medium in the titration vessel is changed for each new sample, the interference of the indication does not take place and a water content can be found reproducibly. Additionally, if the current is reduced from 50 µA to 10 µA, the indication interference occurs later. Since at 10 µA current the titration proceeds much slower, the accelerator is added to the working medium.

- One-component volumetric or coulometric titration
- Sample size: 10 g for volumetry; 5 g for coulometry (needed for reproducibility)



## Applications: Food

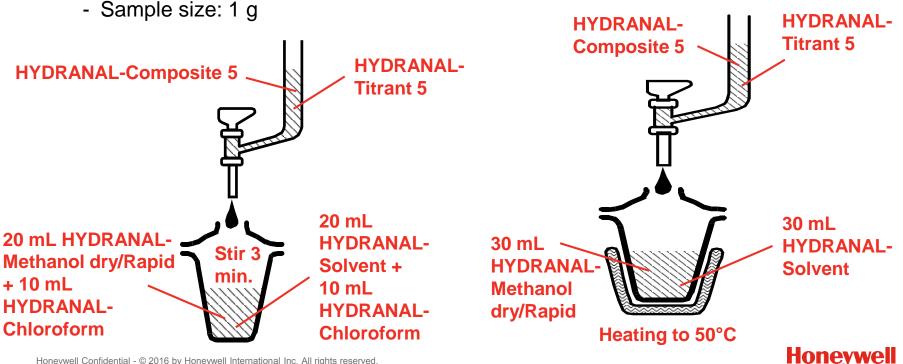


# **Application example: Chocolate (L 071)**

 Water determination in chocolate is trouble-free as long as the sample is dispersed quickly and homogeneously in the solvent. Therefore, methanol is not ideal as solvent. We recommend the addition of **chloroform** in order to dissolve fats. The addition of chloroform can be avoided if the titration is performed at **50°C**. Also, it is advisable to **fractionate** or grind the chocolate.

#### Procedure: •

- One- or two-component volumetric titration

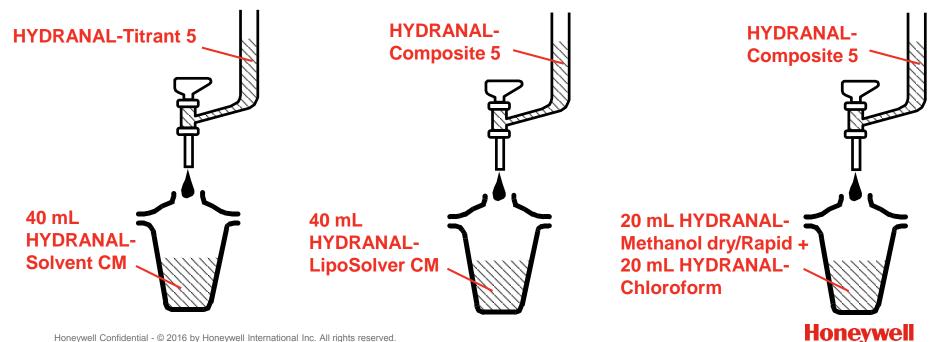


Honeywell Confidential - © 2016 by Honeywell International Inc. All rights reserved.

# **Application example: Butter (L 104)**

• Butter is insoluble in methanol and **chloroform** is a suitable solubilizer. The sample has to be homogenized before analysis. The sample can be introduced directly by means of a **PTFE weighing spoon**, which remains in the solvent until the end of titration. The titration time is about 2 minutes. If a smaller amount of chloroform is used, the dissolving time is longer.

- One- or two-component volumetric titration
- Sample size: 0.5 g



# **Application example: Yeast (L 384)**

- Both fresh and dried yeast did not dissolve in the KF medium and they display differing problems in the extraction of the water.
- Fresh yeast: finely disperses gradually in the titration vessel during the titration. Titration time depends on solvent system used.

#### Procedure:

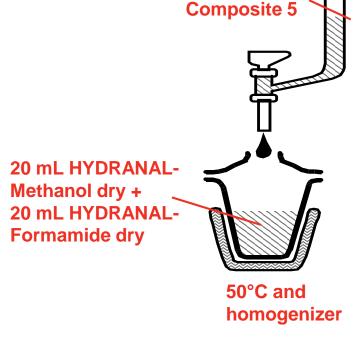
- One- or two-component volumetric titration
- Sample size: 50-100 mg

	Two-component volumetric titration		One-component volumetric titration		
Titrating agent	Hydranal- Titrant 5	Hydranal- Titrant 5 E	Hydranal- Composite 5	Hydranal- Composite 5	Hydranal- Composite 5
Working medium	30 mL Hydranal- Solvent	30 mL Hydranal- Solvent E	30 mL Hydranal- Methanol Dry	30 mL Hydranal- CompoSolver E	20 mL Hydranal- CompoSolver E + 10 mL Hydranal- Formamide dry
Titration time	1.5 min.	2.1 min.	5.3 min.	2 min.	< 2 min.

# **Application example: Yeast (L 384)**

 Dried yeast: the surface of the fine granules is very hard, but should not be mechanically milled in the open air because it is extremely hygroscopic. Dry yeast releases its water content considerably more slowly and the determination times with the above reagents are up to 20 minutes under standard conditions.

- One-component volumetric titration at 50°C with homogenizer and addition of formamide → titration time 2 min.
- External extraction:
  40 g Hydranal-Methanol dry; 1.5 h
- Titration with KF oven: 120°C, titration time 11 min.
- Sample size: 2 g



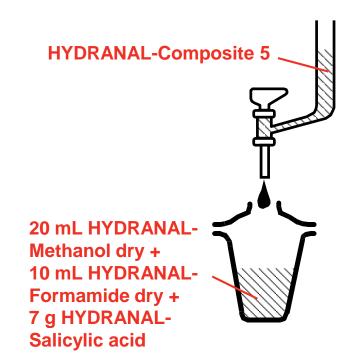


# **Application example: Instant Coffee (L 092)**

- Three types of coffee: freeze dried, spray tower dried and agglomerated.
- They should be titrated in the presence of **formamide** to speed-up the water release.
- In each case to some degree a side-reaction takes place (oxidation by iodine, which is pH dependant). It can be suppressed by lowering the pH value of the working medium conditions with the addition of salicylic acid (to pH 3.2).

#### Procedure:

- One-component volumetric titration
- Sample size: 0.5 g

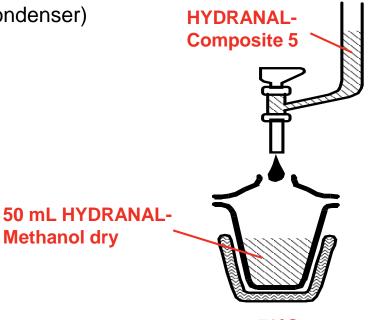


# Application example: Ground Roast Coffee (L 178)

- It is difficult to extract water from natural products, like roast coffee, due to strong cellular bonds. Even at 50°C, the water contained in the coffee could not be satisfactorily extracted.
- Titration in **boiling methanol** is the method of choice for natural products.

### Procedure:

- One-component volumetric titration in boiling methanol (with reflux condenser)
- Sample size: 1 g



**70°C** 

## Closing: Literature and Support

