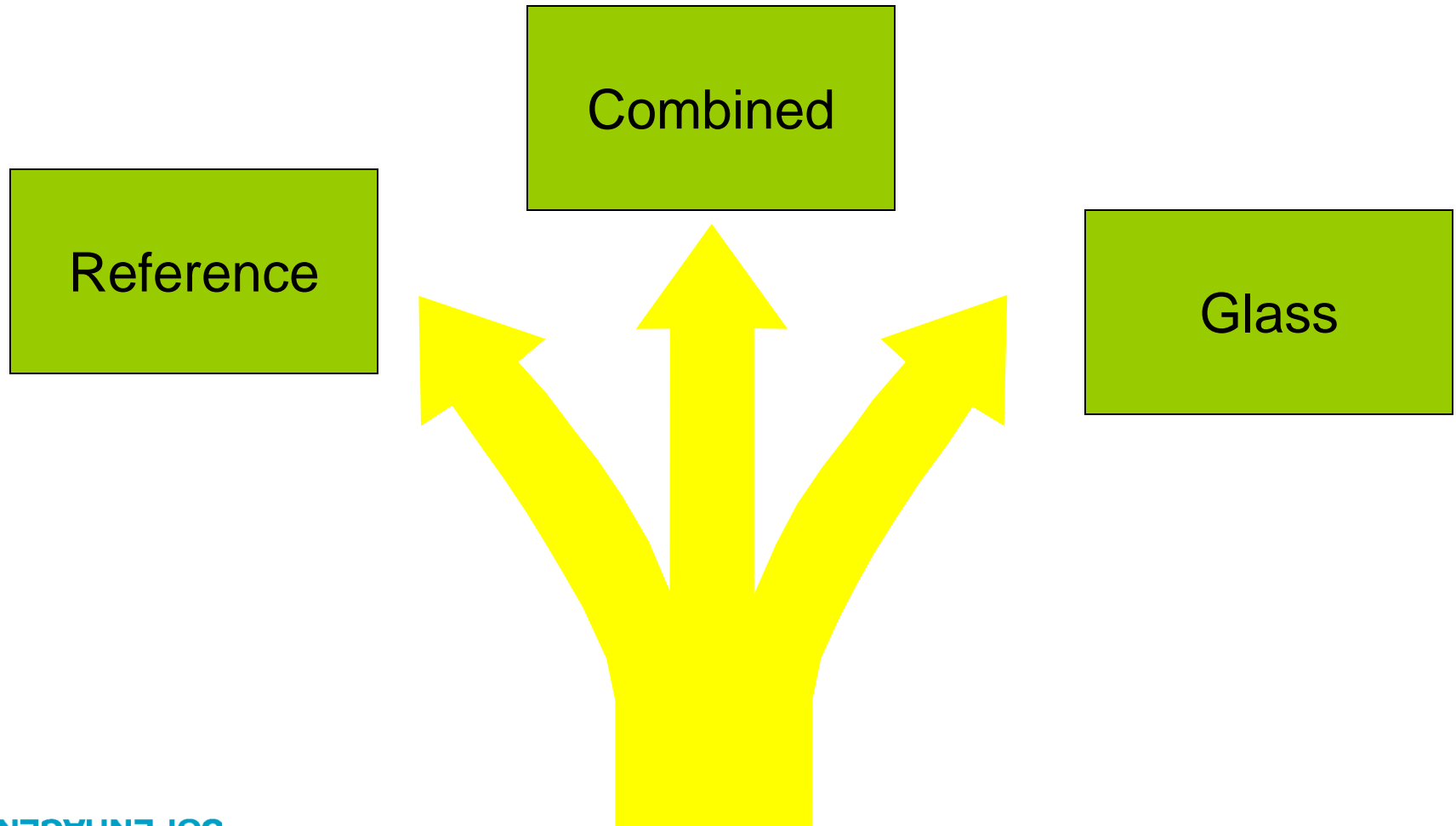


# Electrodes for pH measurements

# Different types



# Electrode working principle

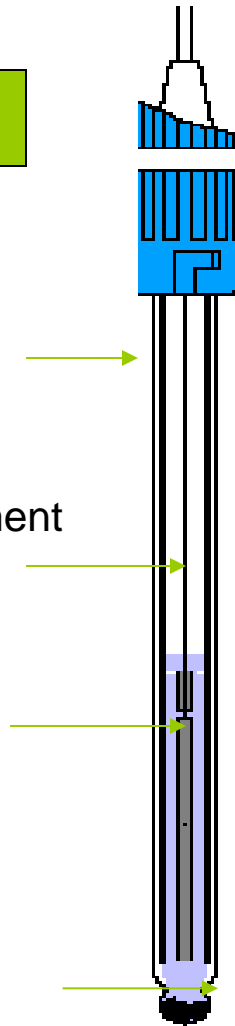
## Glass electrode

Electrode body

Internal reference element

Internal solution

Glass membrane



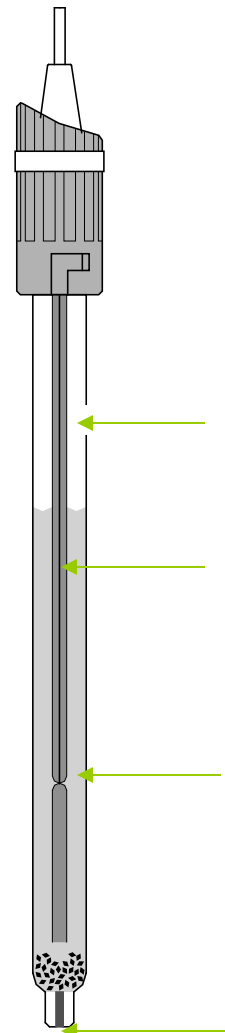
## Reference electrode

Filling hole

Reference element

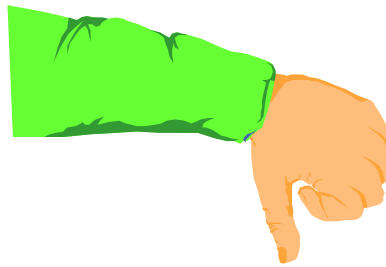
Reference electrolyte

Liquid junction



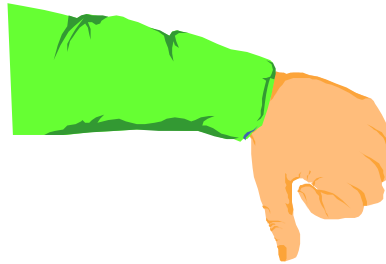
# Sensitive parts

Reference electrode



Liquid junction

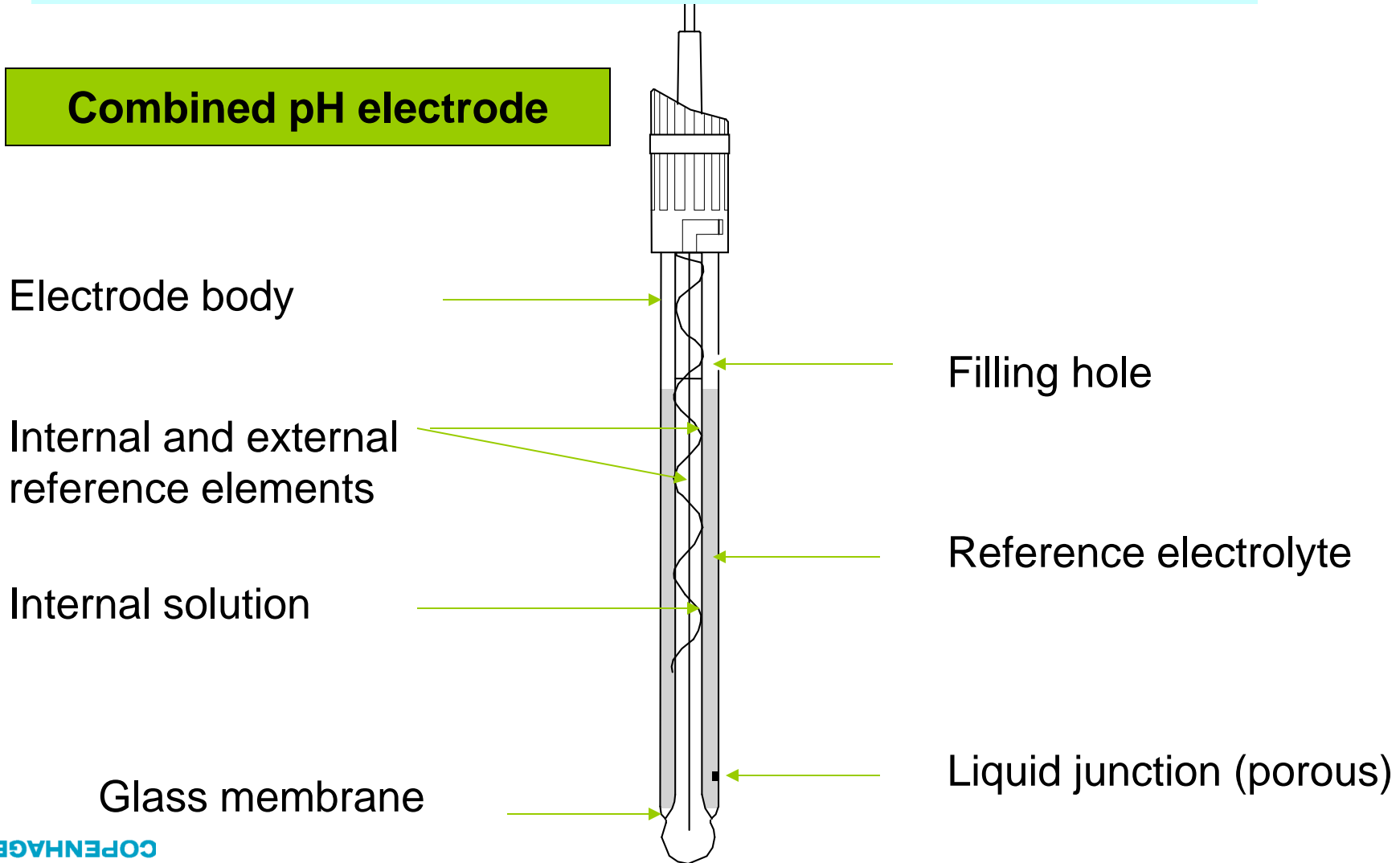
Glass electrode



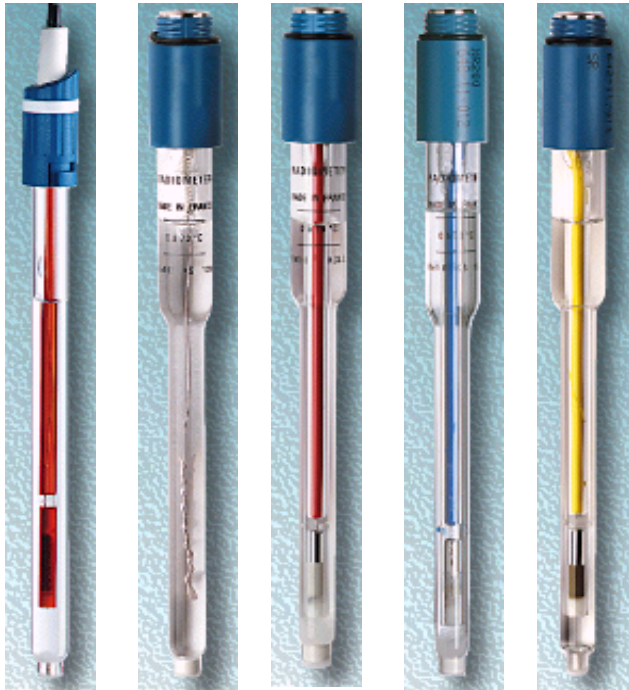
Glass membrane

# Principle of a combined electrode

## Combined pH electrode



# Various reference elements



- Red Rod from Radiometer Analytical
- Silver-silver/chloride ( $\text{Ag}/\text{AgCl}$ )
- Calomel ( $\text{Hg}/\text{Hg}_2\text{Cl}_2$ )
- Mercurous sulphate ( $\text{Hg}/\text{Hg}_2\text{SO}_4$ )
- Mercuric oxide ( $\text{Hg}/\text{HgO}$ )

# The liquid junctions

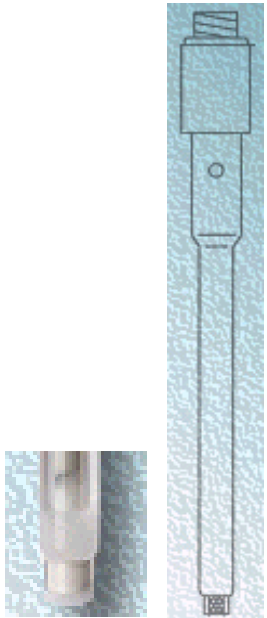
Porous pin

Annular

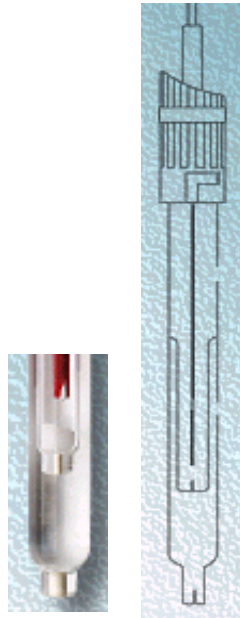
Fibre

Double junction

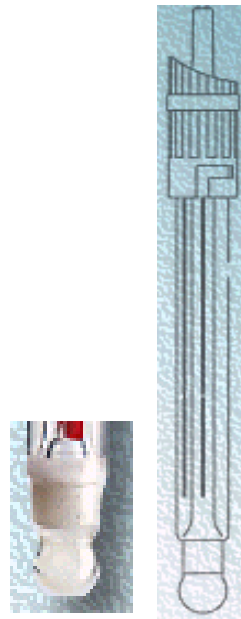
Reversed sleeve



10  $\mu\text{l/h}$



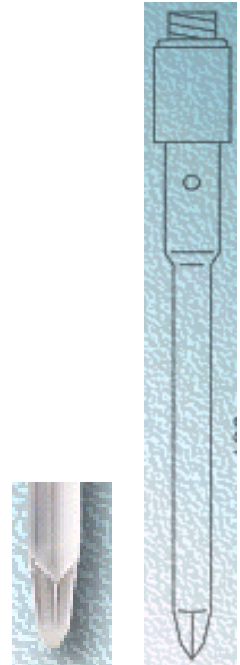
10  $\mu\text{l/h}$



100  $\mu\text{l/h}$



1 ml/h



<10  $\mu\text{l/h}$

# How to select electrodes

Combined electrodes 

Easy to use

Separate reference electrode 

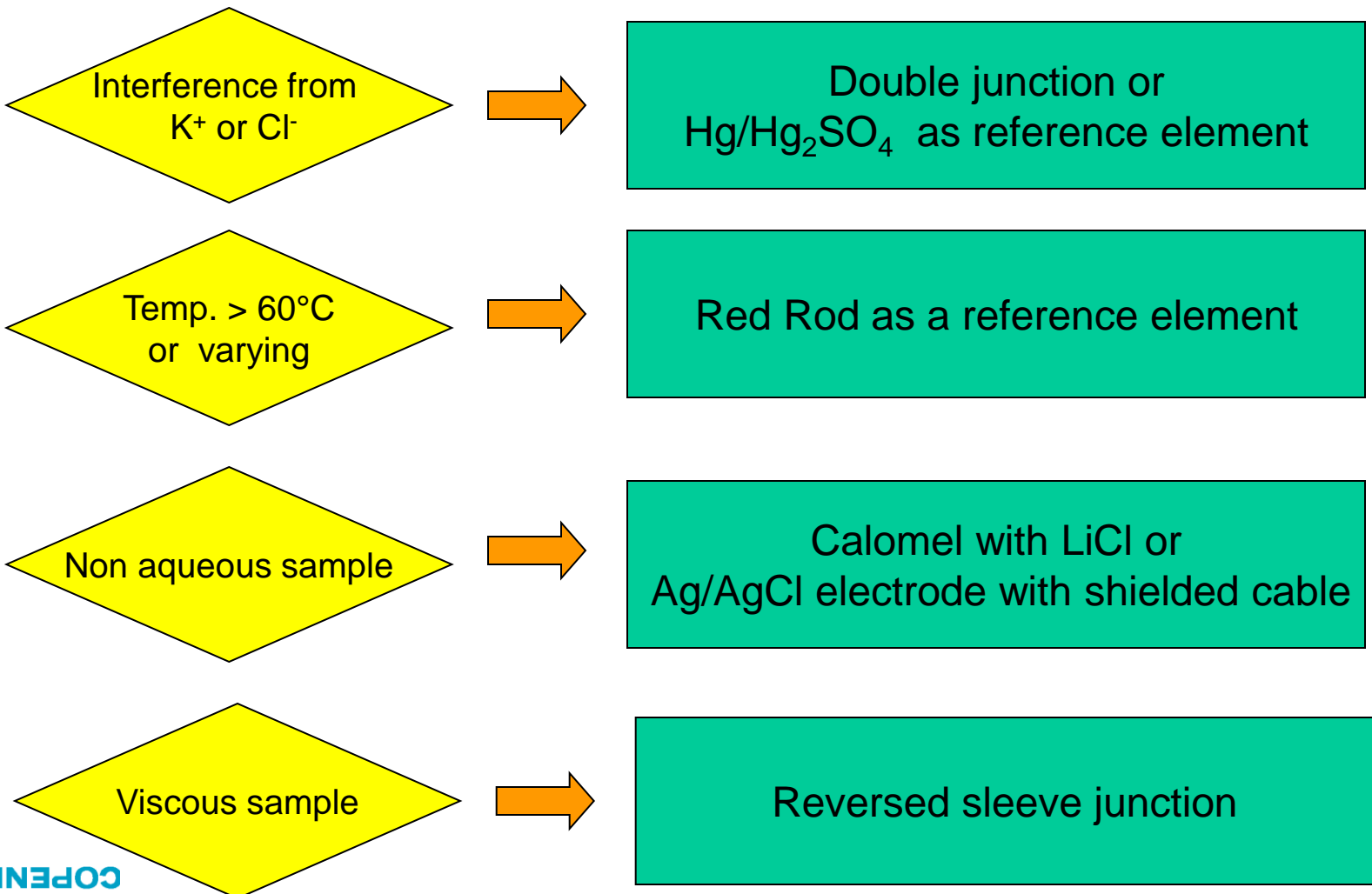
- when liquid junction gets easily clogged

- Non-aqueous solutions or with low ionic strength.

- Electrolyte includes incompatible ions



# How to select a reference electrode



# Red Rod: an innovation

Red Rod combined  
pH electrode

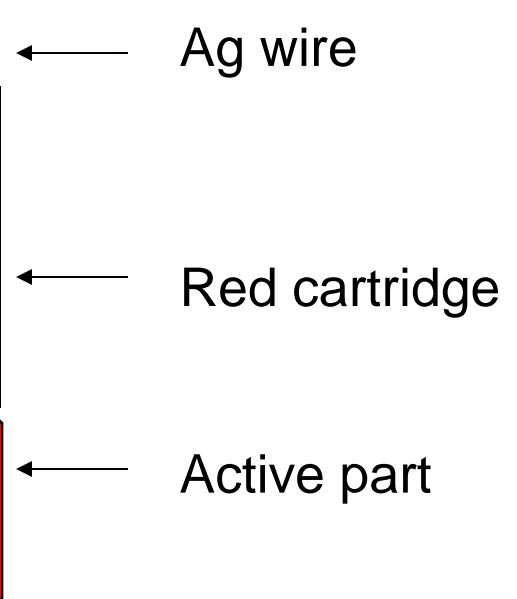
Internal and external  
Red Rod elements

External solution  
(saturated KCl)

Internal solution  
(pH = 6.65)

KCl crystals

Sketch of a Red Rod



# Benefits of Red Rod technology

# Longer lifetime



Less risk of porous pin clogging,  
no **Ag<sup>+</sup> ions** in the filling solution

# Better measurement accuracy



The reference elements are protected from light by the **red cartridge** which means a more stable potential

# Faster response



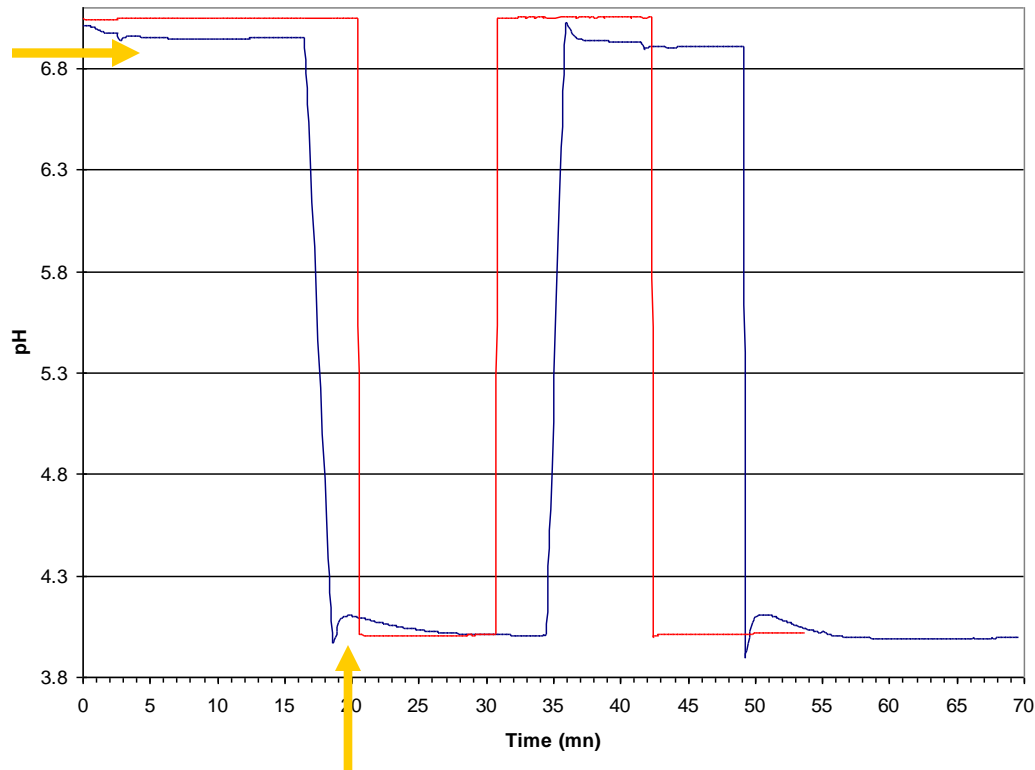
The internal and external reference elements are perfectly symmetrical.

Therefore the behaviour remains the same even when there are large pH or temperature variations.

Measurements from 0 to 100°C

# Reproducible results

Comparison between a Red Rod and a «conventional» electrode



pH 7.000 at 50°C  
pH 4.005 at 25°C

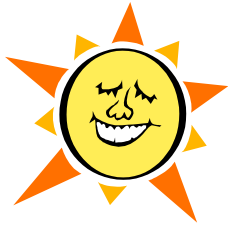
— Traditional Technology  
— 'Red Rod' Technology

# Electrode maintenance

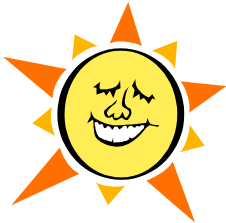


# Electrode maintenance

Correct and regular maintenance means:



Fast response



Reproducible and reliable measurements



A longer lifetime

# Electrode maintenance

## Filling solutions

Type	Description
KS100	saturated KCl, 500 ml
KCl.L	saturated KCl, 100 ml
KCl.C	KCl crystals, 15 g
KCl.Ag	KCl 3M saturated with AgCl, 100 ml
KS110	KCl 3M, 500 ml
KS120	saturated KCl saturated with AgCl, 500 ml
KS160	saturated K <sub>2</sub> SO <sub>4</sub> , 500 ml

## Cleaning solutions

Type	Description
RENOVO.N	Normal cleaning solution, 250 ml
RENOVO.X	Strong cleaning solution, 250 ml
KS400	Pepsine solution in HCl, 250 ml
KS410	Thiourea solution, 250 ml

# Check the state of the electrode



## Reference electrode

- Level of solution (0.5 to 1 cm)
- Crystals:
  - ☐ Present
  - ☐ Move freely
  - ☐ No air bubble

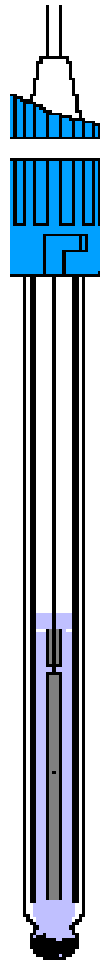
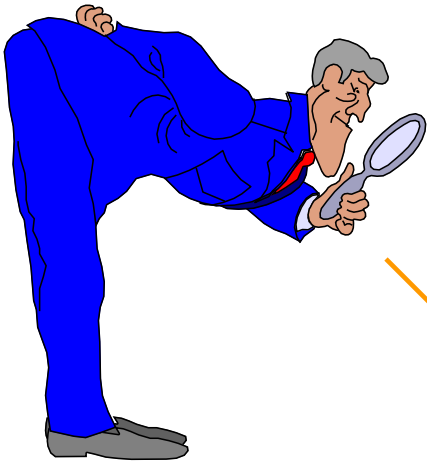
Every day

# Check the state of the electrode

Glass electrode

- Glass membrane:
  - ☐ Clean
  - ☐ No cracks or scratches

Every day



# How to clean

Protein contamination	Immerse electrode in <b>RENOVO.X</b> solution for approx. 3 min. Rinse with deionised water.
	Immerse electrode in <b>KS400</b> solution (pepsin in <b>HCl 0.1 M</b> ) for several hours and rinse with deionised water.
Grease or oil deposit	Rinse with a <b>solvent</b> miscible with water (i.e. <b>acetone</b> ), then rinse with deionised water.
	Close filling hole and immerse in <b>RENOVO.N</b> for 12 to 16 hours.
Silver or sulphide contamination	Close the filling hole and immerse electrode in a <b>KS410</b> ( <b>Thiourea</b> ) solution for several hours. Renew the filling solution.

# GK ANNEX



A complete **kit** for Red Rod electrodes:

- cleaning solutions
- filling solutions
- accessories
- logbook
- instructions

# Troubleshooting

Low sensitivity and/or slow response

Drifting potential

Unstable readings

# Low sensitivity/slow response

Air bubbles

Remove bubbles:

- Tap the electrode with fingers
- Dip the tip in water at 60°C

pH sensitive glass bulb  
contaminated or not hydrated

Identify the contamination

- Use suitable cleaning solution

Hydration:

- At least 1 hour in pH 4



# Low sensitivity/slow response

Calibration error

Buffers:

- Check date of validity
- Use fresh, quality buffers

Electrode too old

Replace the electrode

# Drifting potential

Air bubbles

Remove bubbles:

- Tap the electrode with finger
- Dip the tip in water at 60°C

Level of salt-bridge solution  
too low

Refill with new solution up  
to 0.5 cm under the filling  
hole  
(check there are crystals)

# Drifting potential

Inadequate flow of  
salt-bridge solution

- Check filling hole is uncovered
- If liquid junction contaminated, clean it

No stirring

Reproducible stirring  
during calibration and  
measurements

# Unstable readings

Air bubbles

Remove bubbles:

- Tap the electrode with finger
- Dip the tip in water at 60°C

Liquid junction clogged with crystals from salt-bridge

Replace the filling solution  
(heat the electrode in a  
60°C water bath for 10  
min. to dissolve crystals)

# Unstable readings

Contamination  
with proteins

Lower the electrode tip into:

- RENOVO.X (3 min)
- KS400, pepsin solution (2 hours)

Contamination  
with oil or grease

Lower the electrode tip into:

- RENOVO.N (12 hours)
- Solvent miscible with water (e.g. acetone)

# Unstable readings

Contamination of liquid junction with silver or sulphides

Lower the electrode tip into: KS410, thiourea solution (for a few hours)

Poor connection

Check that the electrode is properly connected and that the meter used is grounded

**The end**