

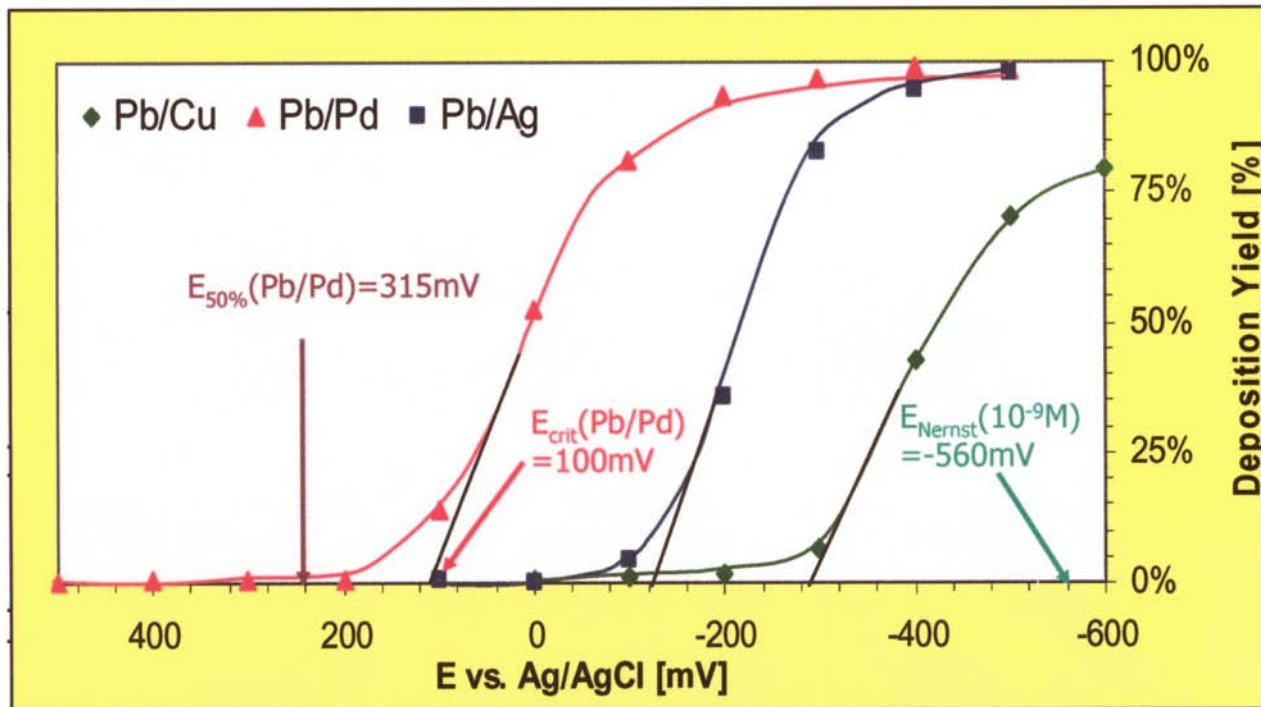
***Electrodeposition Experiments planned
@ TASCA***

J.V. Kratz

Institut für Kernchemie, Universität Mainz

H. Hummrich

Determination of Critical Potentials



Deposition of ^{212}Pb on Pd, Ag and Cu from 0.1 M HClO_4 . Electrolyte Volume $V=1\text{mL}$, Electrode Area $A=1\text{cm}^2$, stirring at 600rpm. The tangents show the E_{crit} values.

The Velocity of Deposition

$$\frac{dN_{\text{dep}}}{dt} = \frac{DF_E}{\delta V} (kN_{\text{tot}} - N_{\text{dep}})$$

F. Joliot, J. Chim. Phys. **27**, 119 (1930)

k = maximum fraction deposited, becomes 1 for a sufficiently negative potential

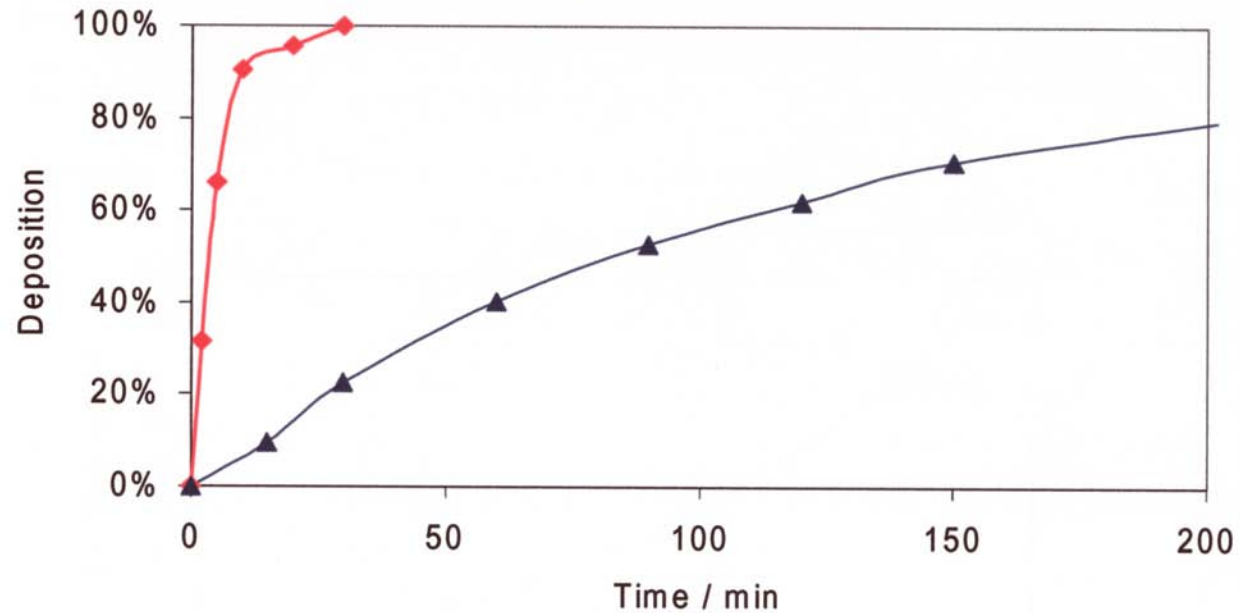
V = volume of electrolyte

FE = area of electrode

D = diffusion coefficient

δ = Nernst diffusion layer

Influence of the V/FE ratio on the Velocity of Deposition

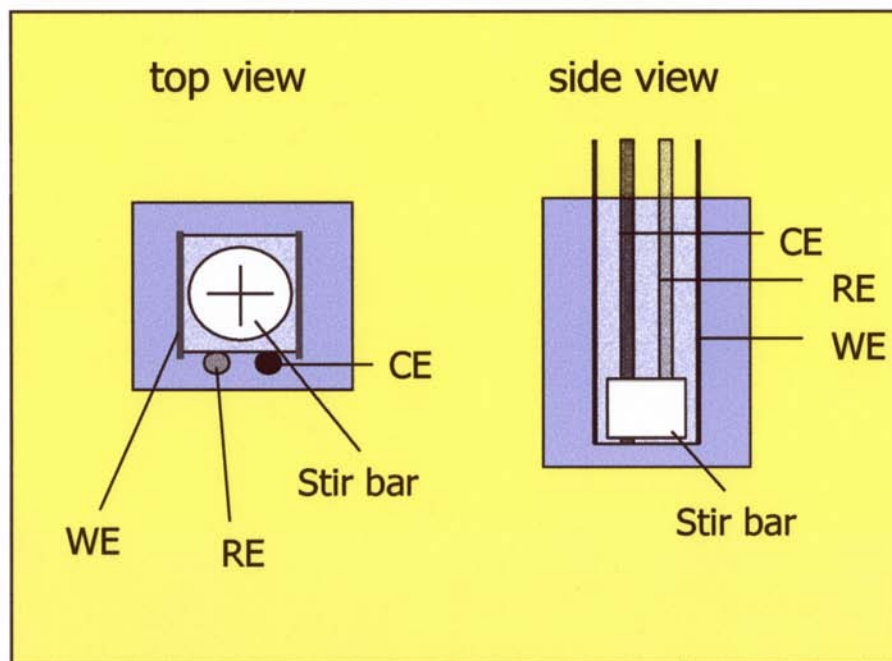


— V = 20 ml, FE = 2 cm², V/FE = 10 ml/cm²

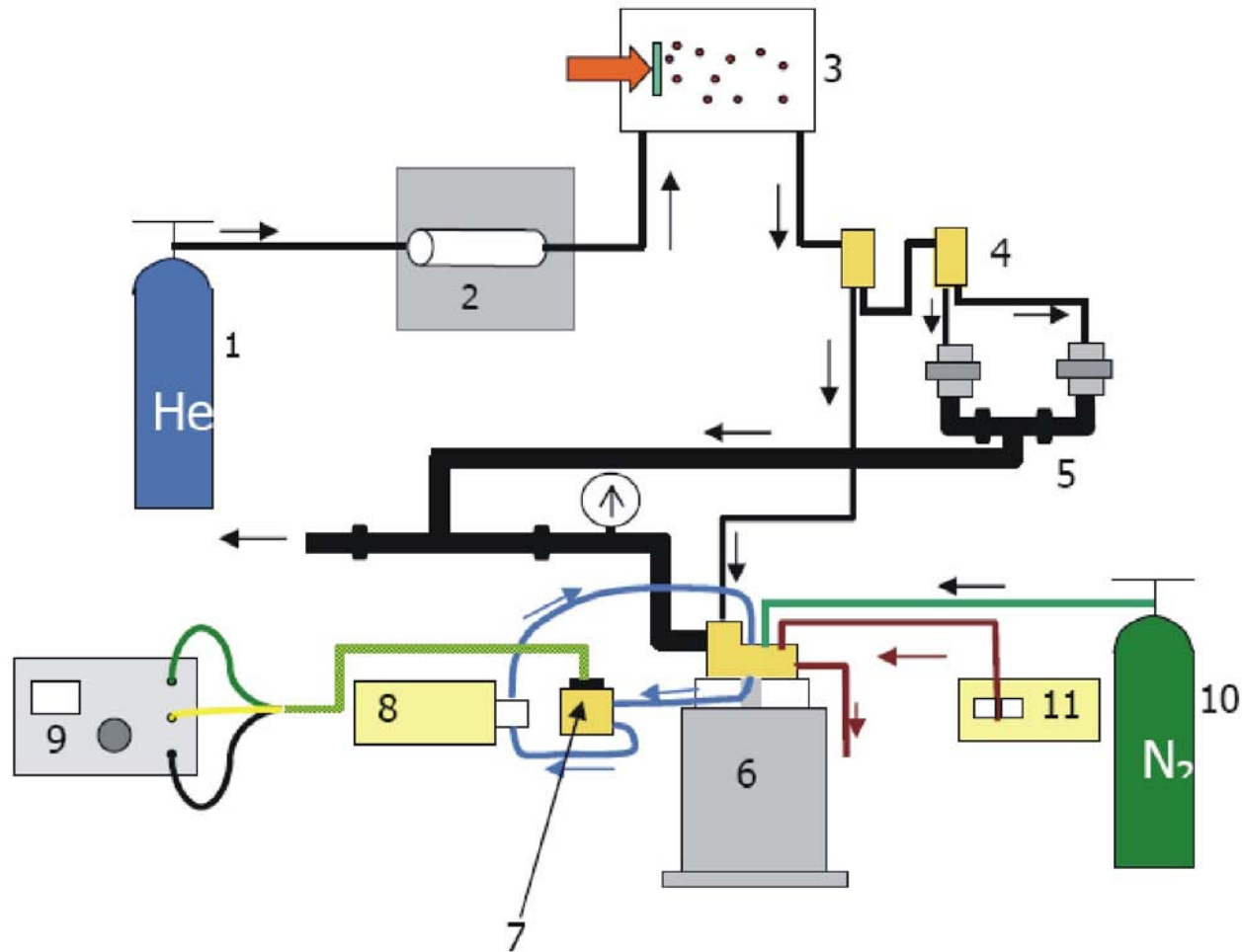
— V = 0,15 ml, FE = 0,36 cm², V/FE = 0,42 ml/cm²

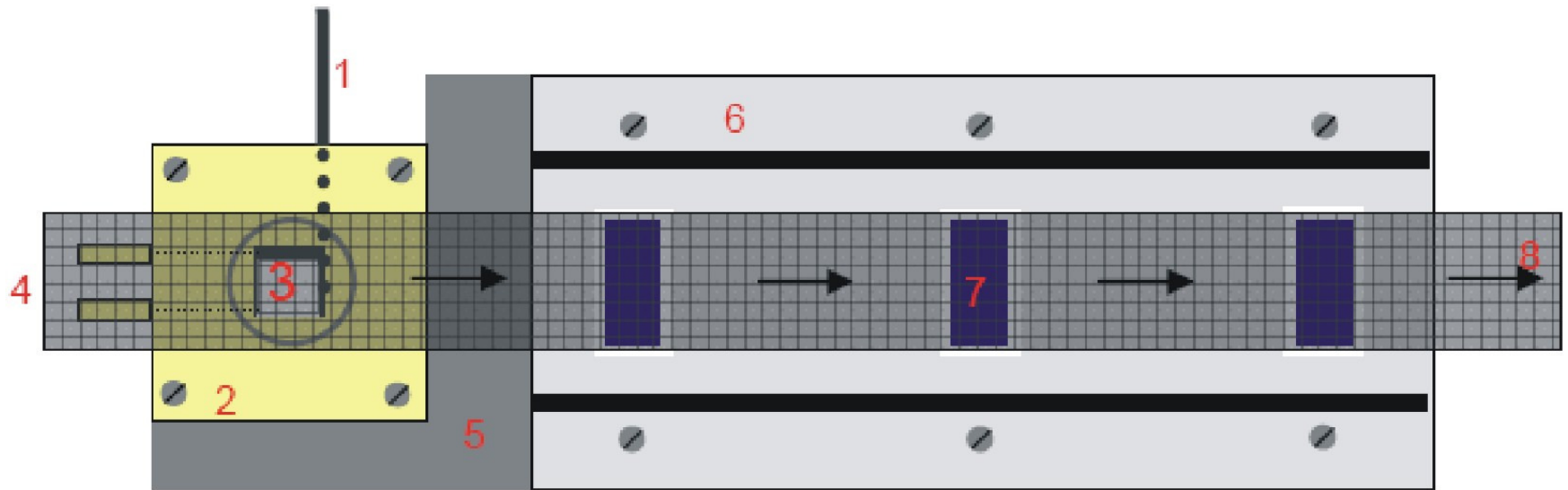
→ Factor 24

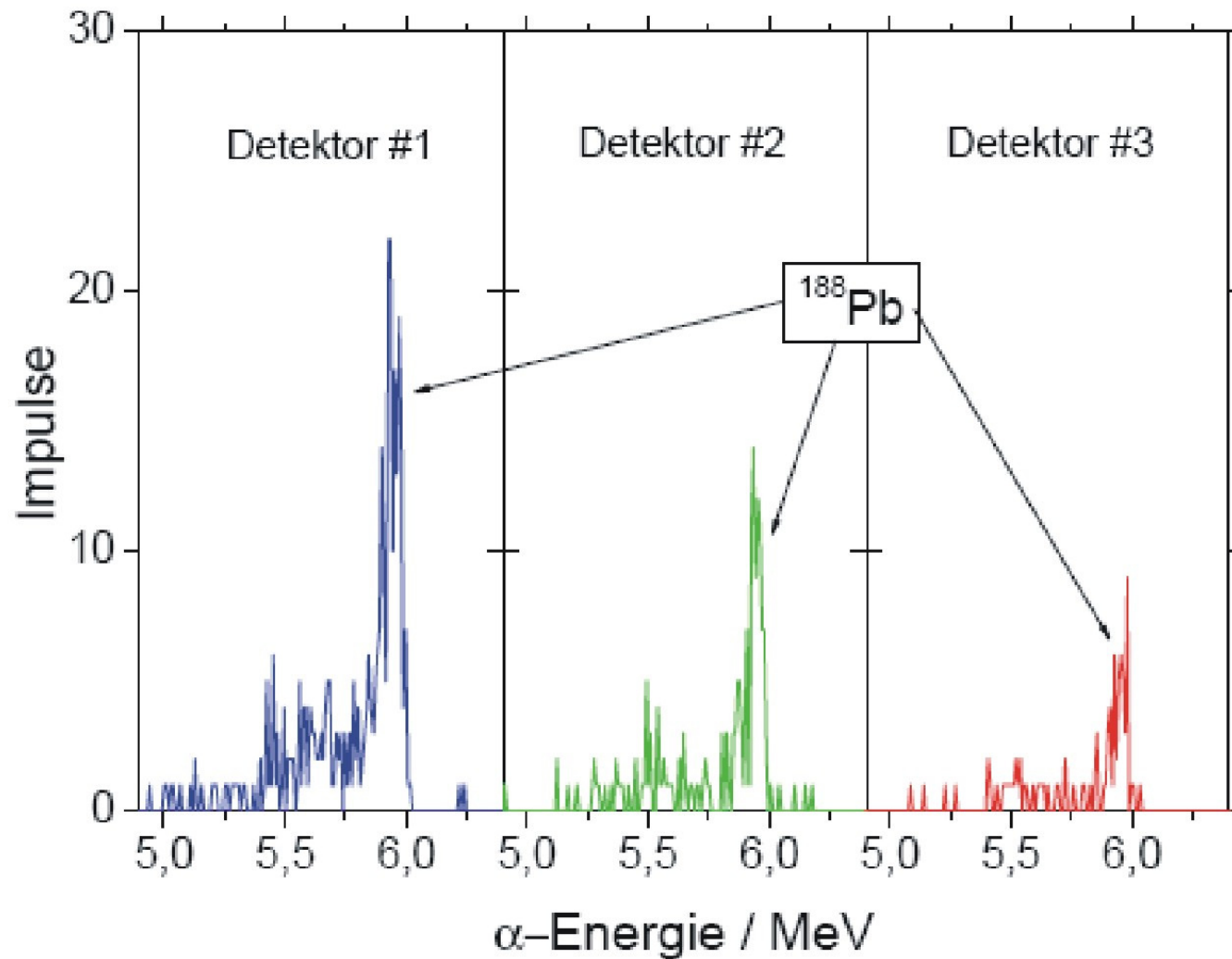
Fast Electrodeposition Cell

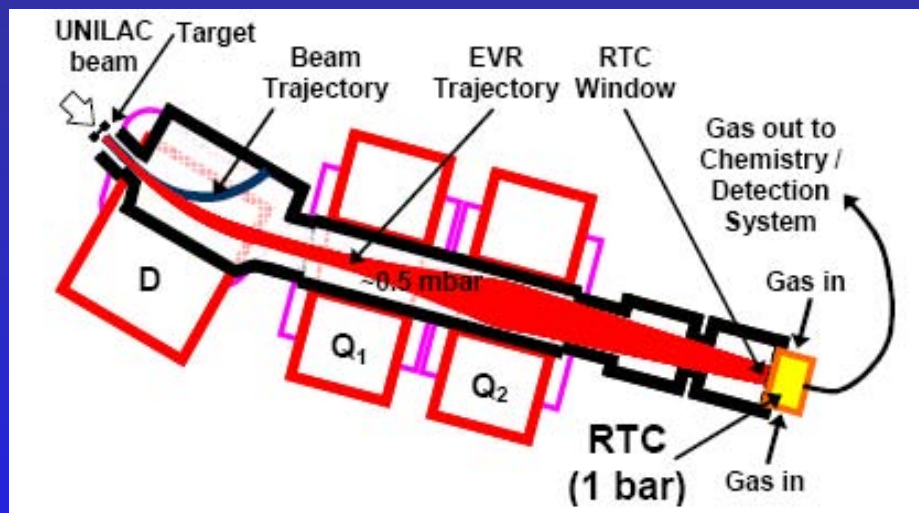


$t_{50\%} = 5\text{s}$ for the deposition of Pb on Pd from 0.1M HClO_4 ,
 $A = 2\text{cm}^2$ and $V = 400\mu\text{l}$ at 90° .
 Stirring with high volume magnetic stirrer at 1400rpm.



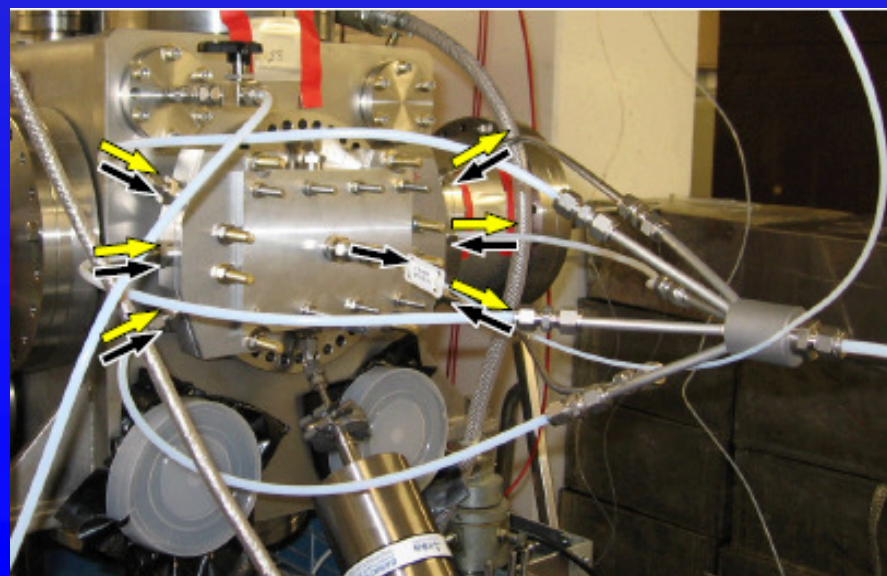


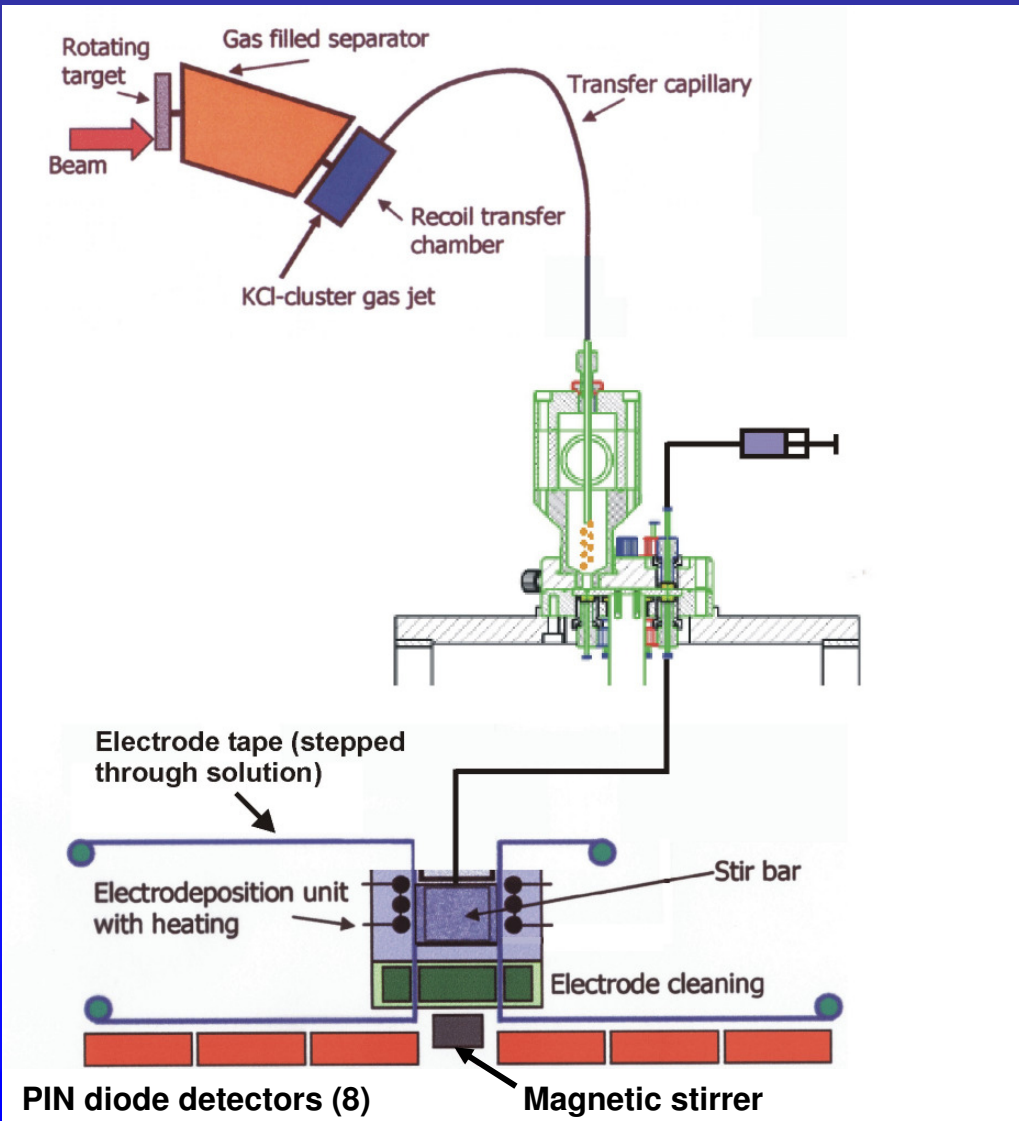




TASCA in the preseparator configuration

The HTM RTC mounted @TASCA.
The two gas – flow regimes are indicated by light and dark arrows.

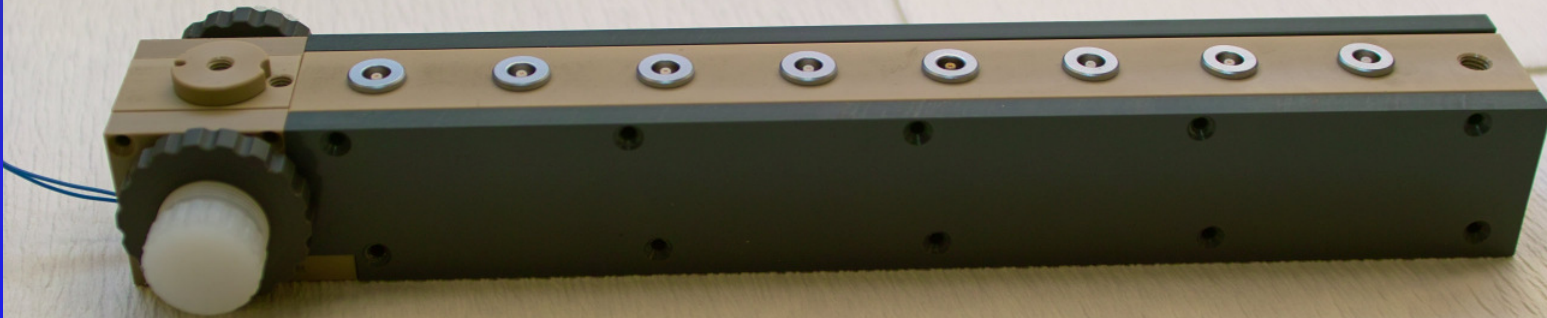




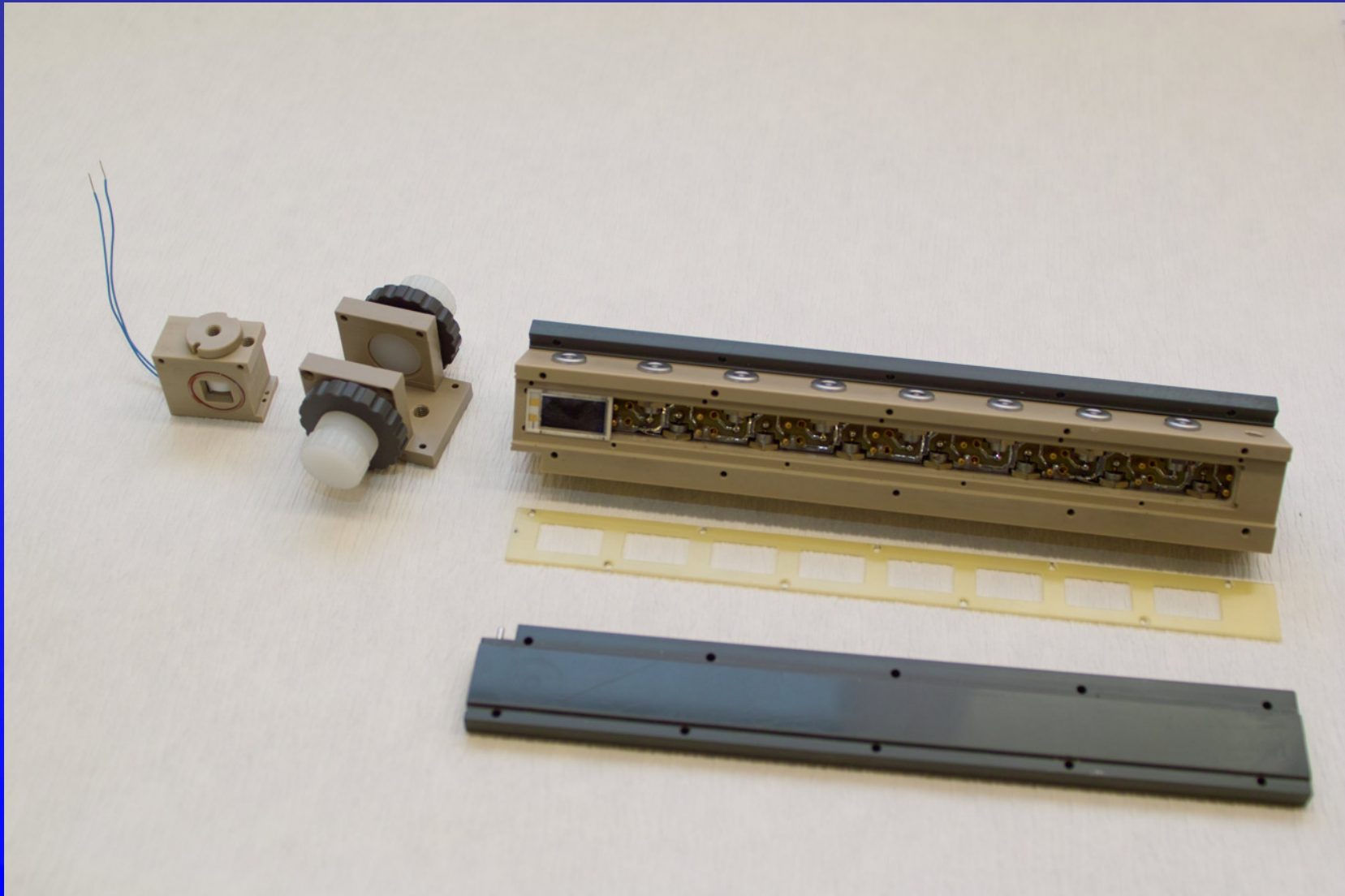
Coupling scheme of

TASCA – RTC – ALOHA –
ELECTRODEPOSITION
with two electrode tapes
and two detector arrays

two detector arrays incorporating
8 photodiodes each



Electrodeposition cell for two
movable tapes





α 8.88 MeV



SF 400 ms