

The Neutrino Mass from Tritium β Decay

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- Introduction
- Tritium β decay experiments in Mainz
- The improved Mainz Experiment
- *Analysis and Discussion*
- Summary and Outlook

Present group members:

Jochem Bonn
Lutz Bornschein
Beate Degen
Christian Weinheimer
E. O.

Visitors: from Troitsk:

Vladimir Lobaschew
Oleg Kazachenko
Juri Jeraskin
⋮

from Dubna/Prague:

Alois Kovalik

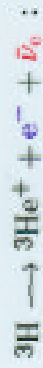
Collaborator in T_2 -film physics:

Paul Leiberer (Konstanz)

Fresh Diploma Students as from 1999:

Lars Fickinger
Christine Kraus
Holger Ullrich

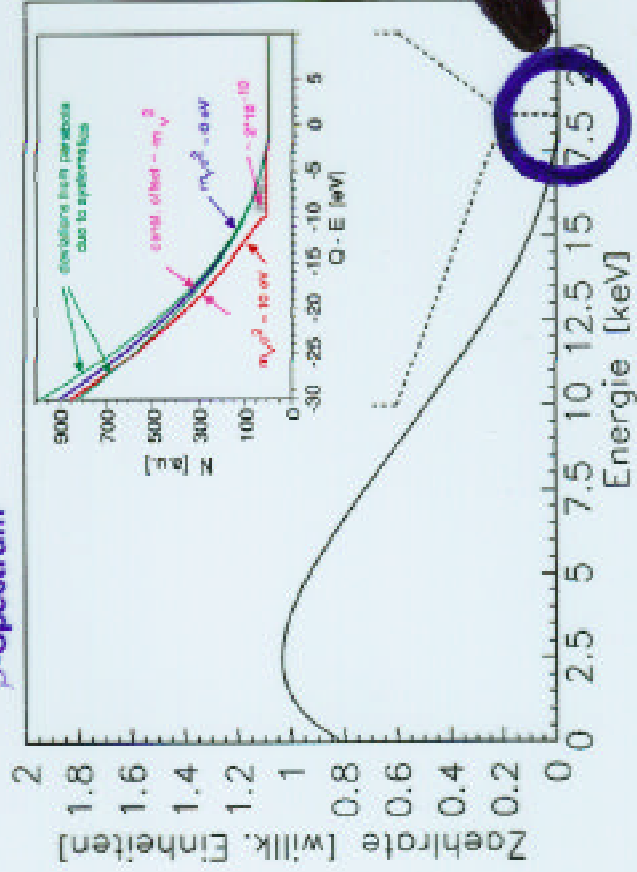
Direct measurement of m_{ν_e}



superallowed transition

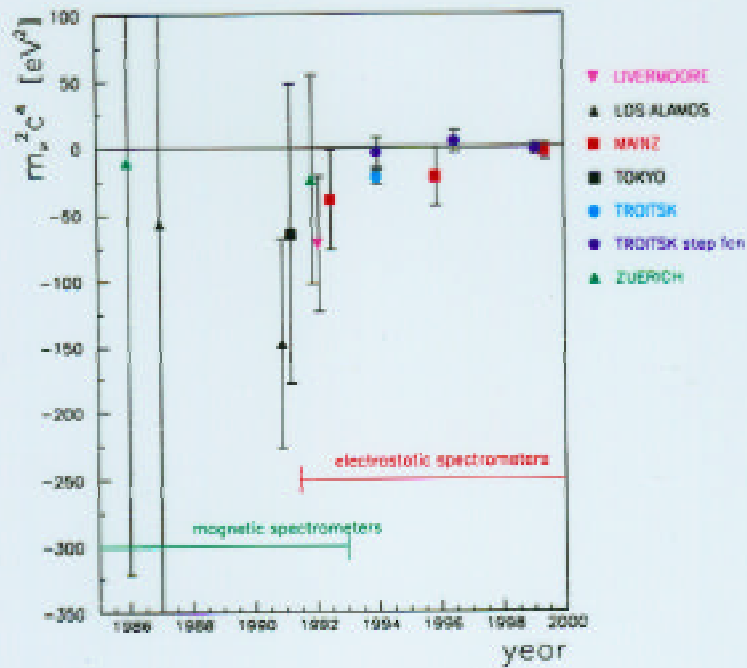
$Q = 18.6 \text{ keV}$, $t_{1/2} : 12.3 \text{ a}$

β -Spectrum



if more than one mass eigenstate contributes (not resolved): $\overline{m}_\nu^2 := \sum_i |U_{ei}|^2 \cdot m_i^2$

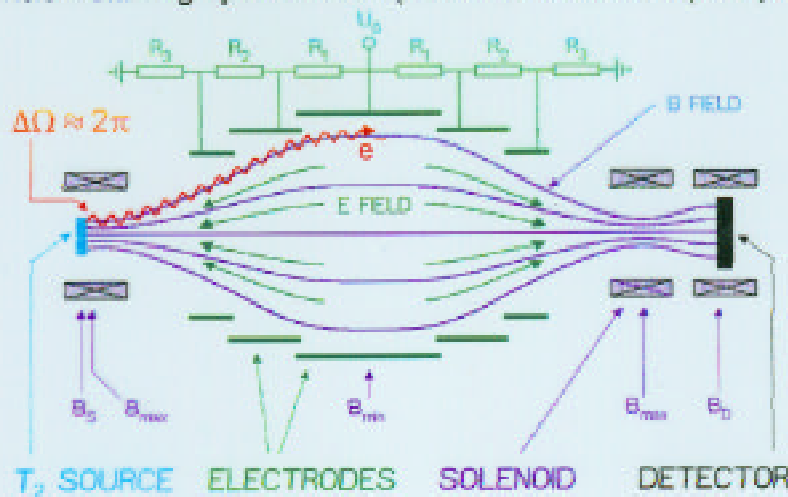
Tritium β decay: last decade



- 2 running experiments: Mainz, Troitsk
- Mainz 1991/1994 and Troitsk 1994:
 - only data close to endpoint: $m_\nu^2 \approx 0$
 - down to 500 eV below endpoint: $m_\nu^2 \approx -100\text{eV}^2$
 - undetected additional energy loss

Magnetic Adiabatic Collimation + Electrostatic filter

"Solenoid Retarding Spectrometer" (Nucl. Inst. Meth. B63 (1992) 345)



$$\vec{F} = (\mu \vec{\nabla}) \vec{B} + q\vec{E} \quad \mu = \frac{E_{\perp}}{B} = \text{constant}$$

WITHOUT E FIELD:



- magnetic guiding field

$$\rightarrow \Delta\Omega \approx 2\pi$$

- adiabatic transf. $E_{\perp} \rightarrow E_{\parallel}$ + electrostatic retardation

$$\rightarrow \Delta E \approx 4 - 6 \text{ eV}$$

- performs point to point image
- keeps off background originating outside flux tube

INTENSITY

MONOENERGETIC ELECTRONS →

ENERGY

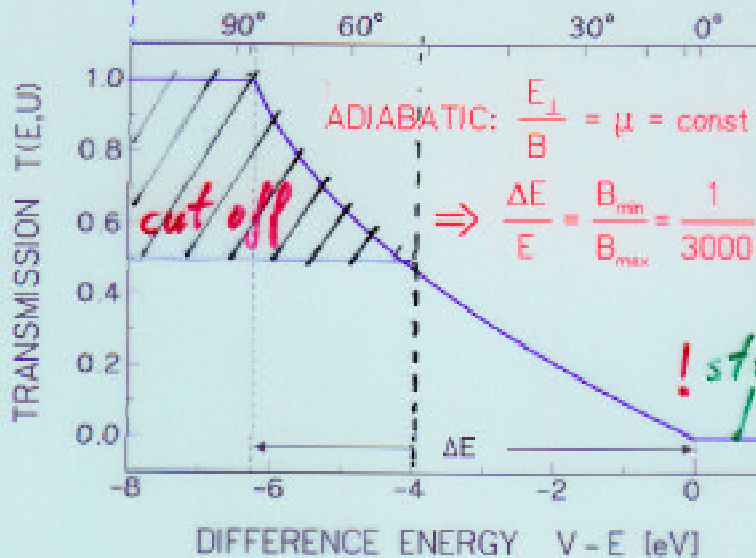
E

COUNT RATE

POTENTIAL V

E

MAX. TRANSMITTED STARTING ANGLE

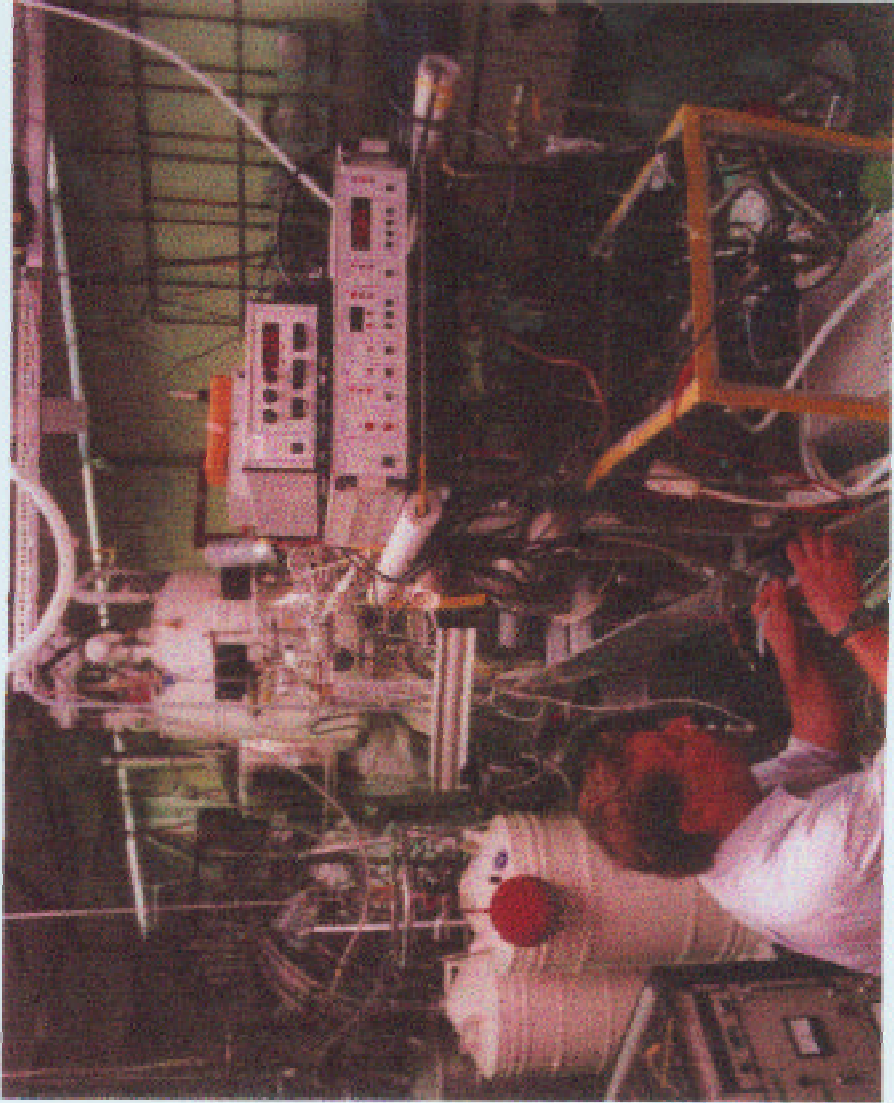


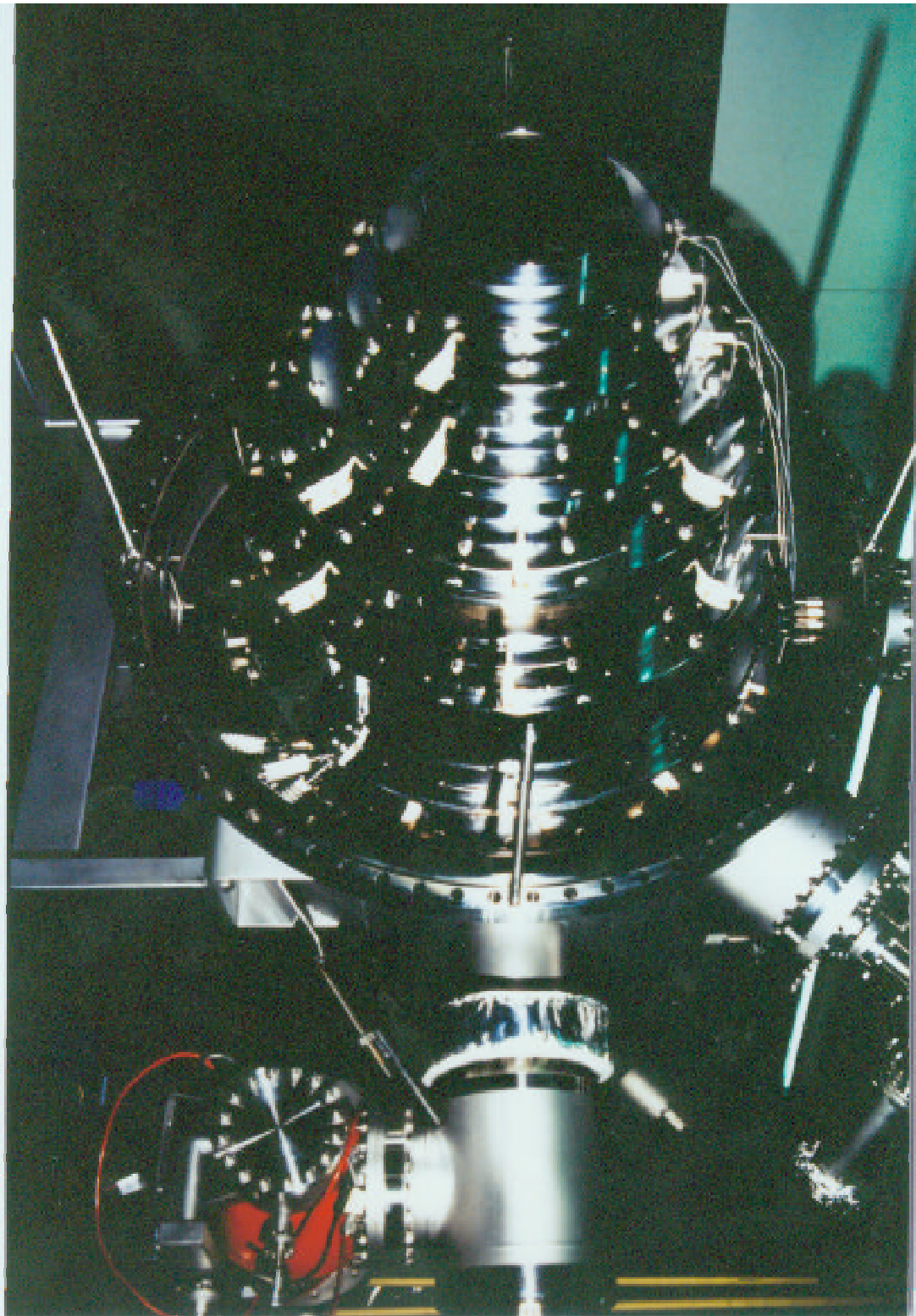
ISOTROPIC SOURCE:

$T(E, V)$

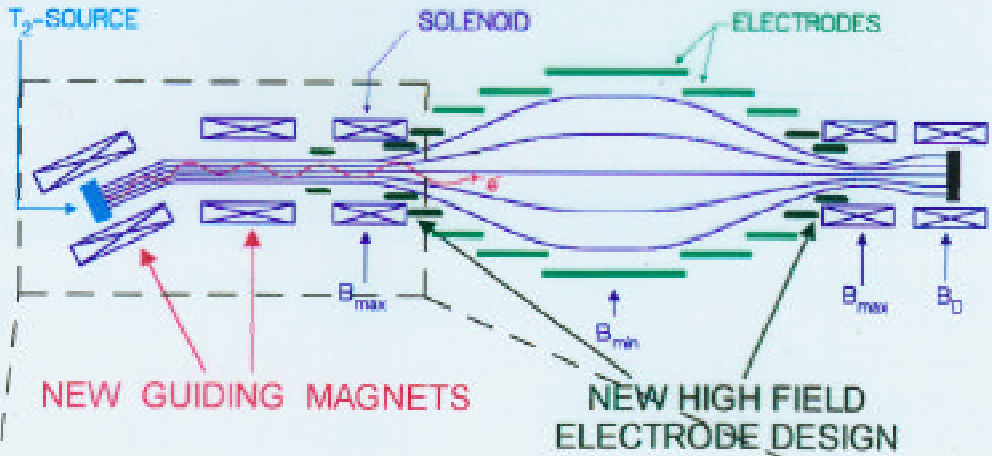
$$T(E, V) = \frac{1 - \sqrt{1 - \frac{(E - V) \cdot B_s}{E \cdot B_{\min}}}}{1 - \sqrt{1 - \frac{B_s}{B_{\max}}}}$$







IMPROVEMENTS OF THE EXPERIMENTAL SETUP



NEW GUIDING MAGNETS

NEW HIGH FIELD ELECTROBE DESIGN

LHe COOLED TRAP

- tritium mol. from source are cryotrapped
- residual gas mol. from spec. are cryotrapped

NEW LHe CRYOSTAT

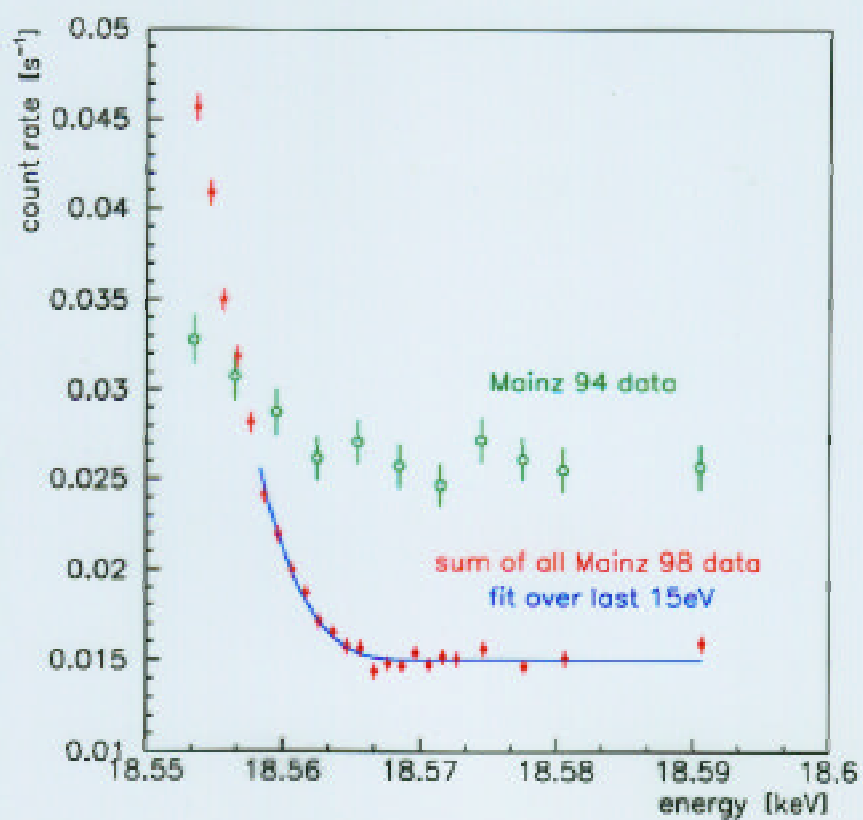
- $T > 1.7\text{ K}$ adjustable
- larger source area $d = 20\text{ mm}$
- max starting angle $78^\circ \rightarrow 45^\circ$
- monitor film roughness

NEW SUPERCONDUCTING GUIDING MAGNETS

- electrons are guided from source to spectrometer
- tritium molecules not!

Summing up all 1998 data

Q3+Q4+Q5



The Fitfunction

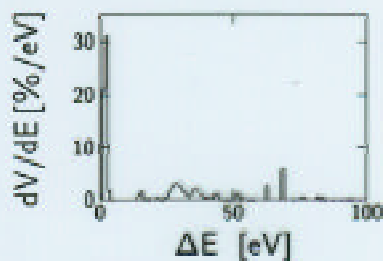
$$N(E) = \text{amp} \sum_i P_i \varepsilon_i^2 \sqrt{1 - \frac{m_{\nu}^2 c^4}{\varepsilon_i^2}} \otimes f_{\text{response}} + \text{BG}$$

$f_{\text{resp}} \approx \text{Charging} \otimes \text{TF} \otimes \text{EL} \otimes \text{BS}$
 with $\varepsilon_i = E_0 - V_i - E$: final states

final states:

sudden approx.

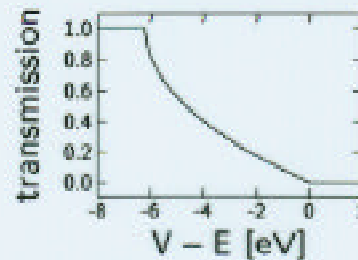
(Kolos, Fackler)



TF:

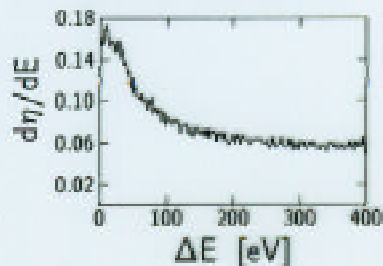
transmission fcn

(analytically known)



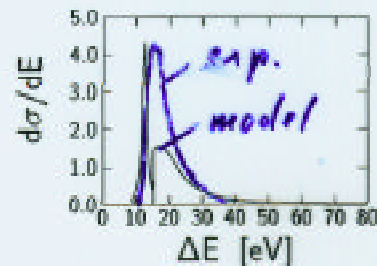
BS: backscatter:

MC-calculations



EL: energy-loss:

$^{83\text{m}}\text{Kr}$ -measurements



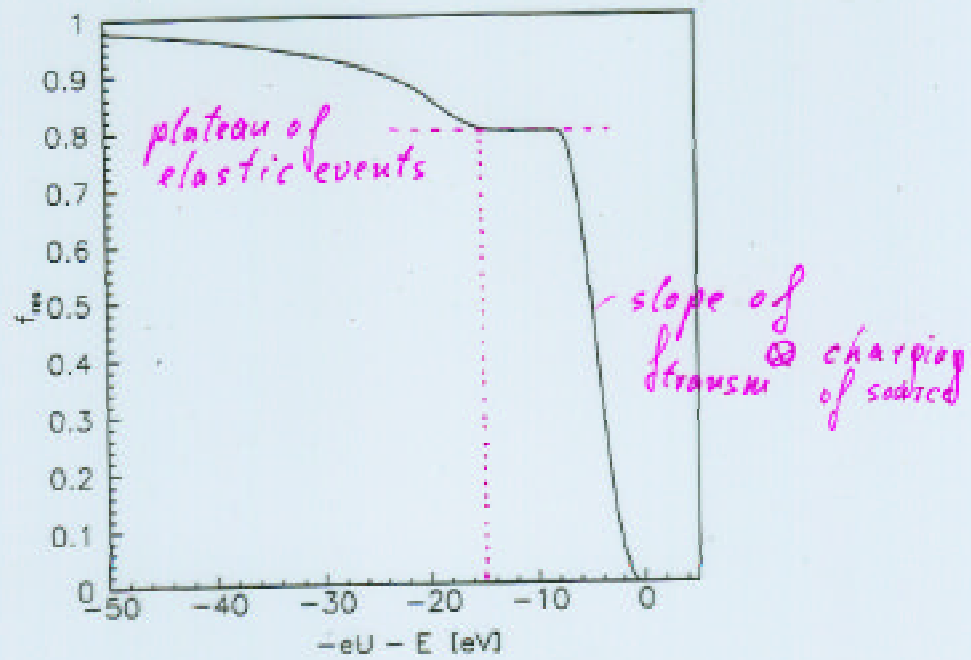


Figure 2. Normalised response function of the spectrometer f_{res} for a monoenergetic electron source of energy E in dependence on the retarding energy $-eU$. The energy loss f_{loss} and the charging effect f_{charge} are calculated for a source thickness of 490 \AA as used for measurements Q3-Q5.

Systematic uncertainties

- **Inelastic Scattering** **film thickness** (49 %)



T_2

λ_{free}

shape of energy loss function

- **Final states** (effects due to solid state) (37 %)

spectator excitation

changes of excited states energy levels

- **T_2 film charging up** **new** (14 %)



- **T_2 film roughness** (avoided) (-)



T_2

- **Backscattering** (small) (< 1%)



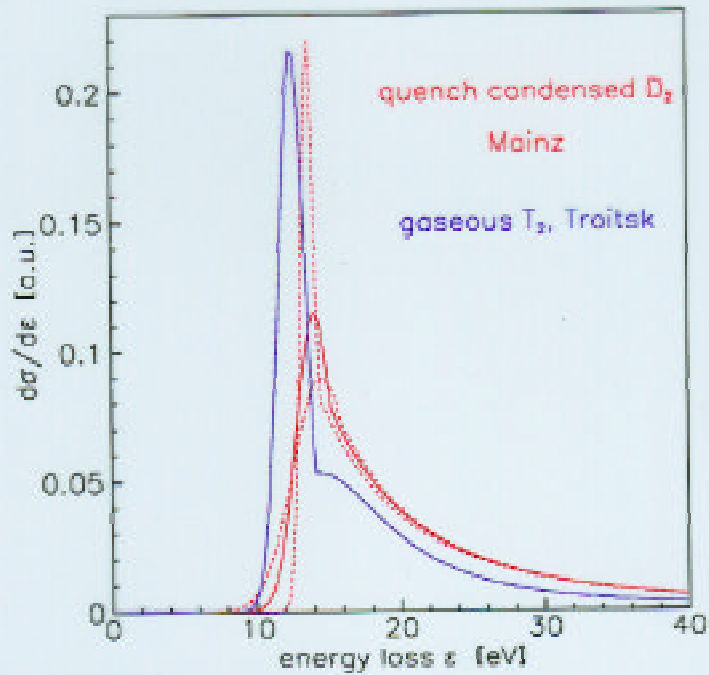
T_2

carbon

- **Detector efficiency** (small) (-)

(contribution to systematic uncertainty for Q5, last 70 eV \leftrightarrow)

Energy loss function



Determination with conversion e^- through D_2 films:

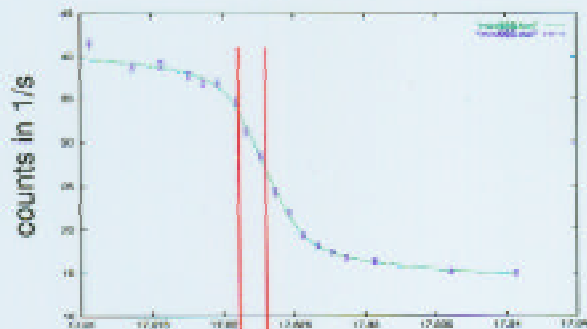
- → shape for quench condensed film is significantly different to shape for gas!
Influence of Pauli blocking for excited molecular states!
- → $\sigma_{\text{tot}} = 3.00 \pm 0.16 \cdot 10^{-18} \text{ cm}^2$;
14% smaller than expected
Influence of Pauli blocking for excited molecular states!

Charging up of T₂ film

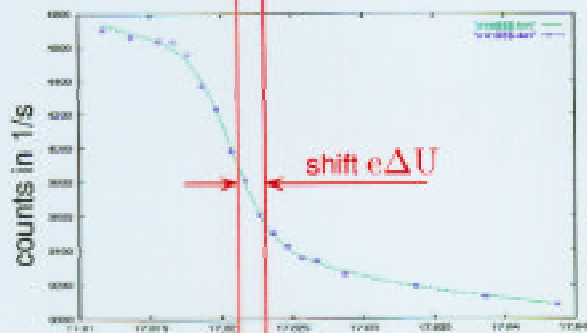
Shift of endpoint energy (run 1997)

Charging up of the T₂ film (40 mCi)

→ Measurements with K conversion electrons of Kr-83



Kr-83 submonolayer
on top of 240 ML D₂

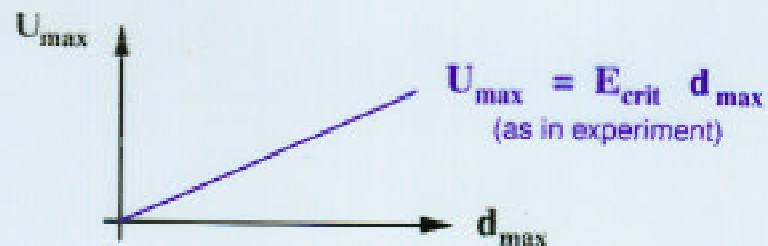


Kr-83 submonolayer
on top of 120 ML T₂

energy in keV

Does microscopic picture fit?

1) Expectation from picture:



2) Measured critical field E_{crit} :

$$E_{\text{crit}} = 6 \text{ V} / 288 \text{ Monolagen}$$

$$\rightarrow W_{\text{pass}} = e \cdot E_{\text{crit}} \cdot d_{\text{ML}} \approx 250 \text{ K}$$

(right order of magnitude !)

⊕ charging up effect probably understood

⊖ deterioration of energy resolution

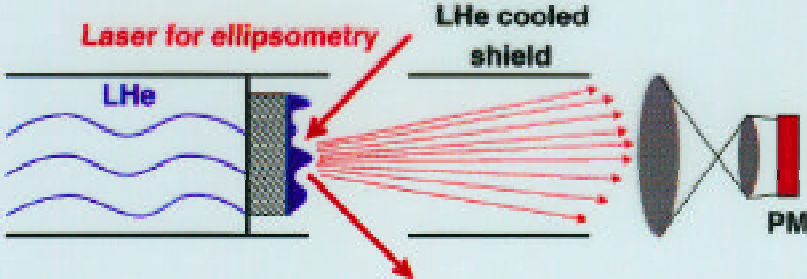
(10% – 90% : 3.5 eV → 6.0 eV)

• take effect into account in analysis

estimate for uncertainty: $\approx 50\%$ of total effect

Detection of Roughening Transition

Observation of stray light

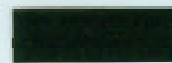
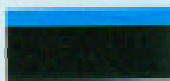


Temperature Dependence of Roughening Transition

condensation

roughening transition

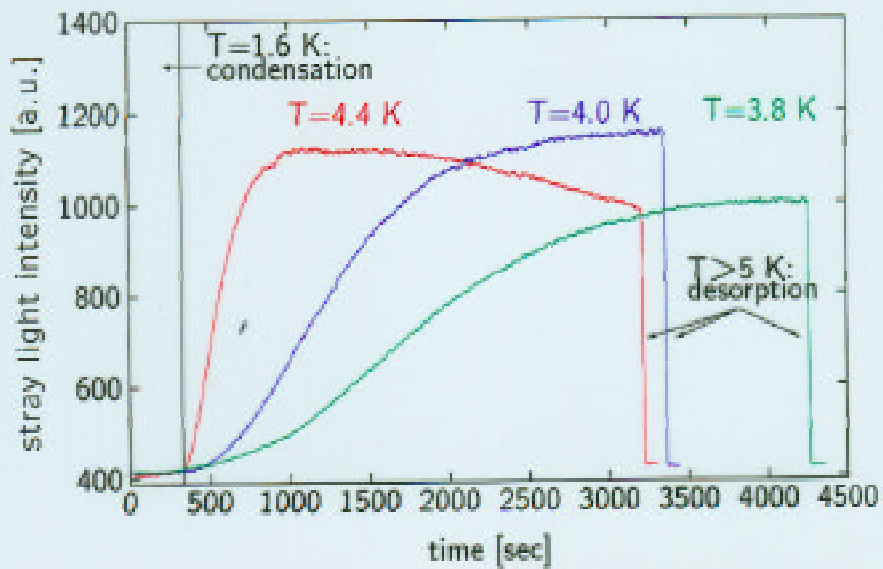
desorption



1.6 K

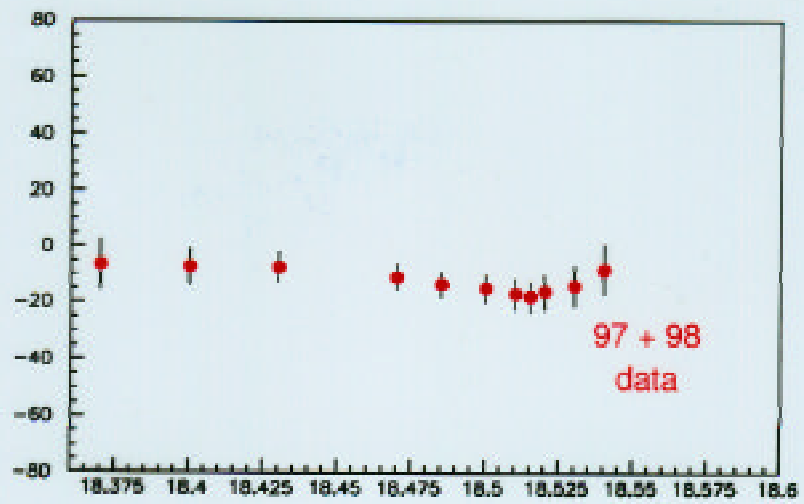
3.8-4.4 K

> 5 K

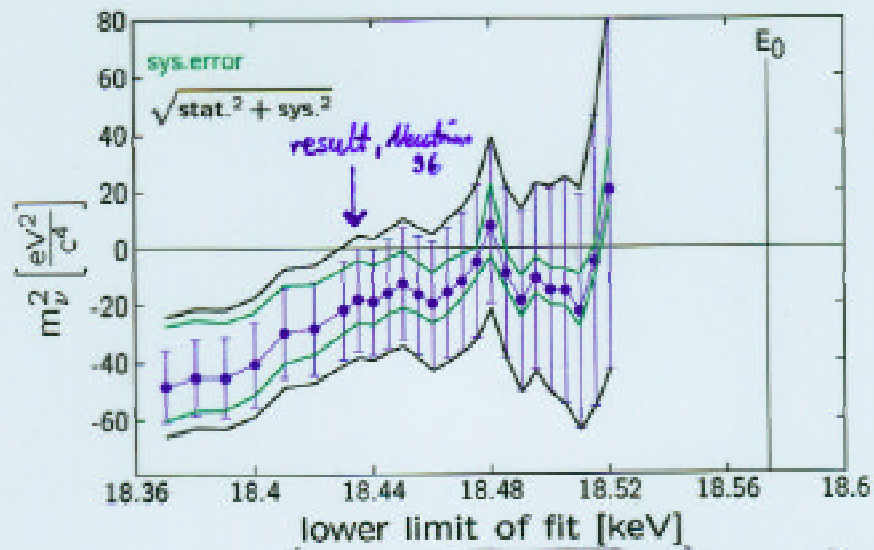


Extrapolated time constants for T_2
(ok, if T_2 decays do not matter)

T	4.2 K	2.8 K	1.6 K
τ	hours	days	years



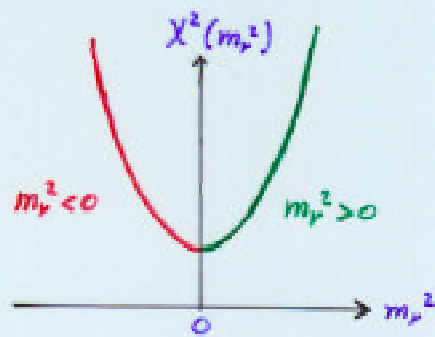
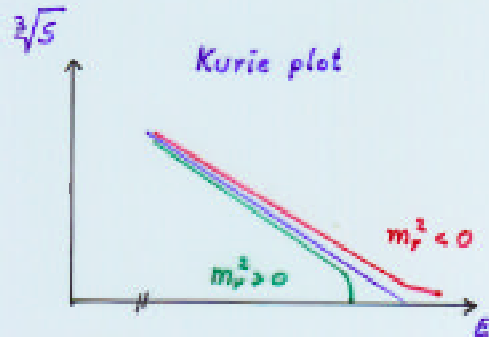
Result 1994



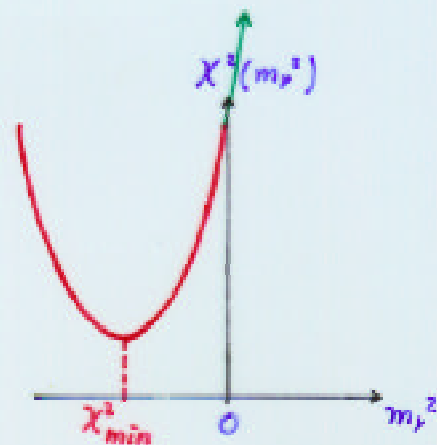
- fit abuses fit parameter m_ν^2 to compensate systematic discrepancies between data and fit function
- trend explainable by missing energy loss

How to treat negative m_ν^2 - values in the fit function ?

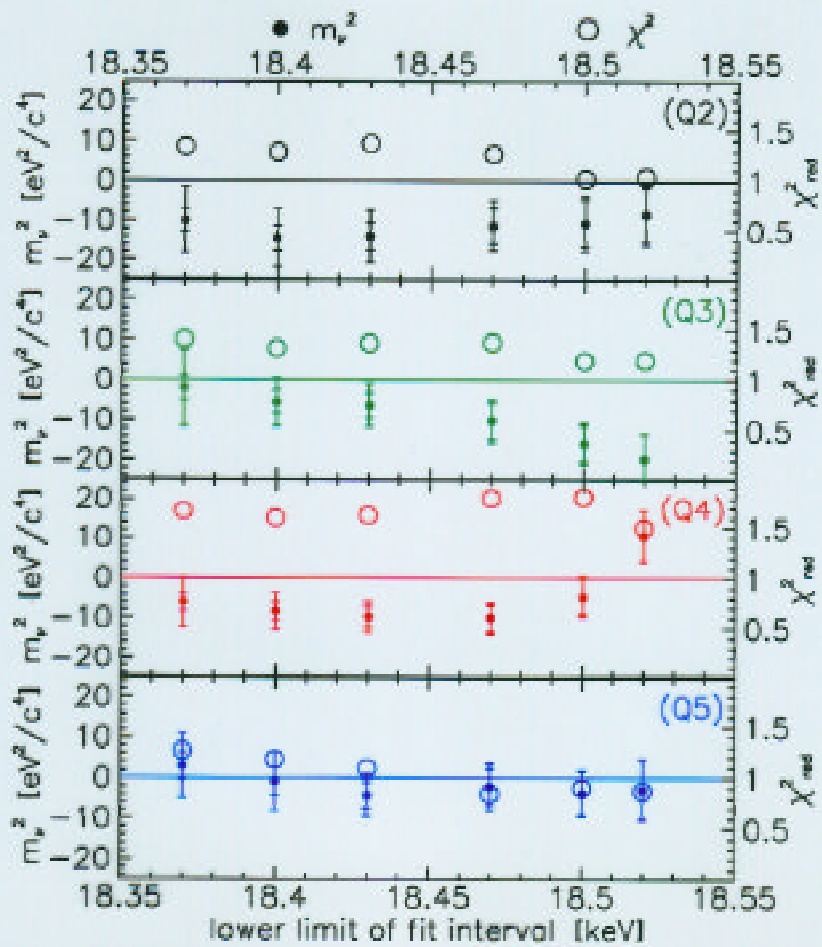
Find a suitable analytical continuation of β -spectrum beyond E_0 which produces a parabolic $\chi^2(m_\nu^2)$ - curve around $m_\nu^2 = 0$ to accommodate statistical fluctuations of excess count rate



However:
systematic distortion of
spectrum, e.g. by under-
estimated fraction of
inelastic processes
produce significant but
unphysical negative
 m_ν^2 -values



$m_\nu^2(\text{fit})$ versus fit interval



Mainz upper limits on neutrino mass 2

2.) If one believes in step as correct solution:

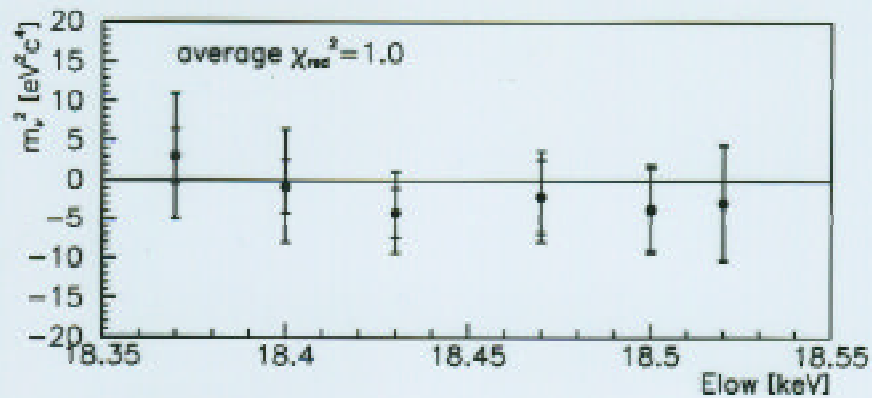
→ fit step and subtract it (only possible for Q4):

last 70 eV (Q4, 1998 run II):

$$m_\nu^2 c^4 = -1.8 \pm 5.1 \pm 2.0 \text{eV}^2$$

$$\rightarrow m_\nu c^2 \leq 3.0 \text{ eV} \quad (95\% \text{C.L., unified approach})$$

3.) Considering Q5, 1998 run III alone:



→ Standard analysis:

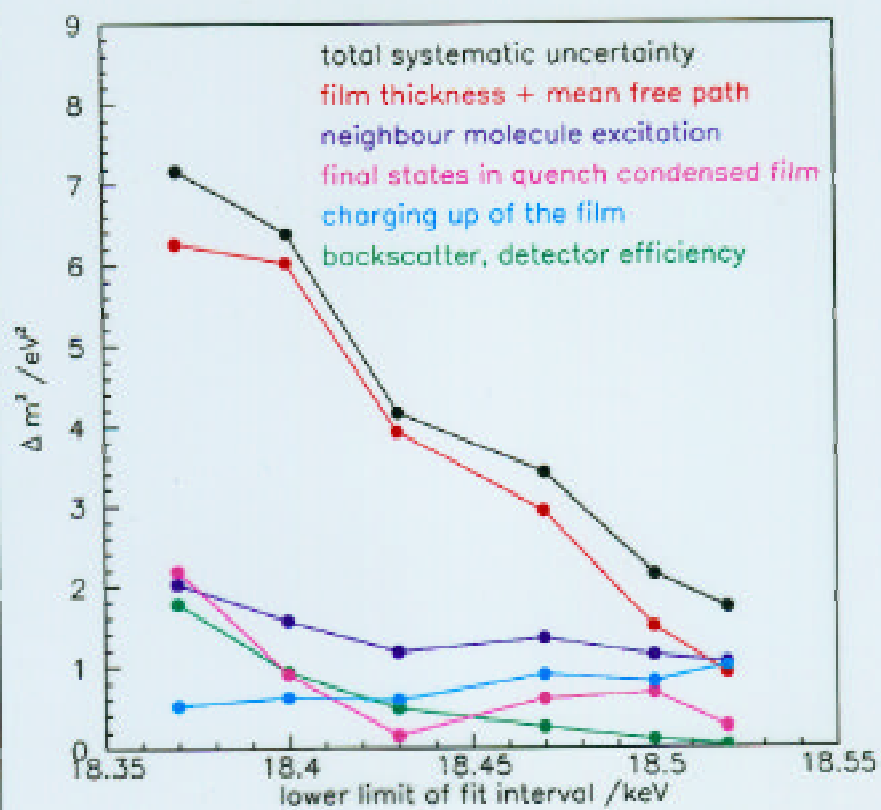
last 70 eV (Q5, 1998 run III):

$$m_\nu^2 c^4 = -3.7 \pm 5.3 \pm 2.1 \text{eV}^2$$

$$\rightarrow m_\nu c^2 \leq 2.8 \text{ eV} \quad (95\% \text{C.L., unified approach})$$

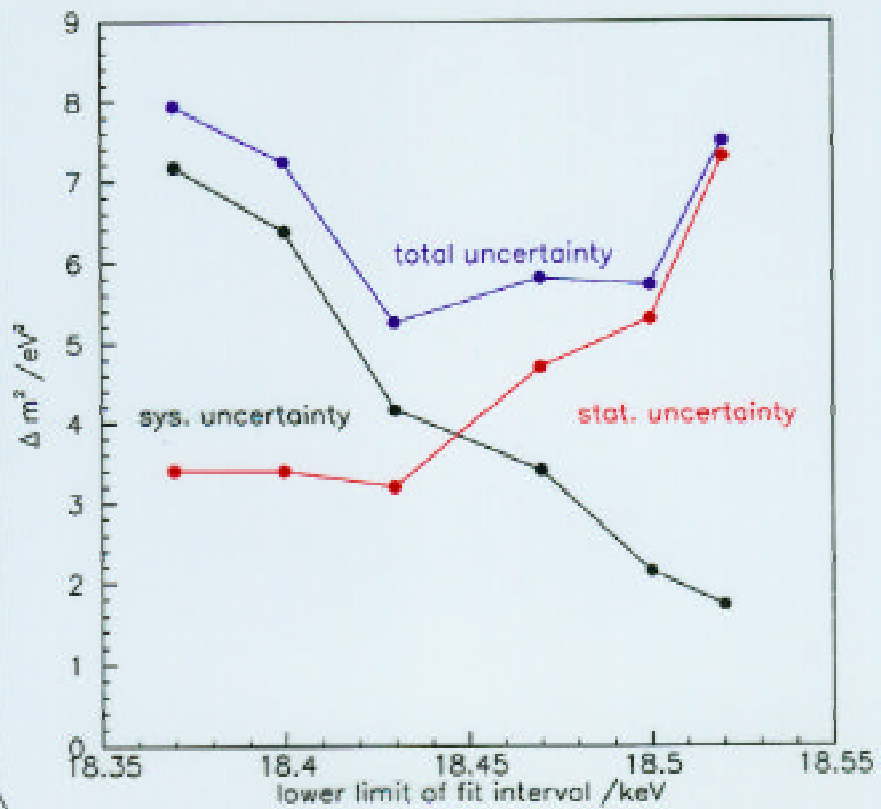
Systematic uncertainties

1998 run III: Q5



Statistical and systematic uncertainties

1998 run III: Q5

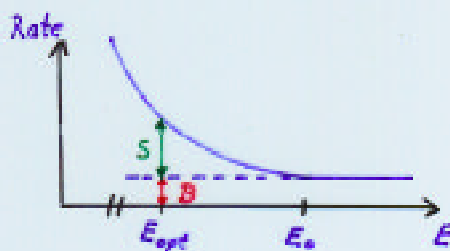


Statistical Error δm_ν^2

given: **S** = Signal rate
B = Background rate

$$\delta m_\nu^2(E) = \frac{2}{3c^4} (E_0 - E)^2 \frac{\sqrt{(S+B)/t}}{S}$$

Optimal point in the spectrum



$$S(E_{\text{opt}}) = 2B$$

There the dependence on

- source strength **R**
- background rate **B**
- measuring time **t**

is given as

$$\delta m_\nu^2 \propto R^{-\frac{2}{3}} \cdot B^{\frac{1}{6}} \cdot t^{-\frac{1}{2}}$$

Correlation of errors

- of $(\text{mass})^2$: δm_ν^2 and
- Endpoint E_0 : δE_0

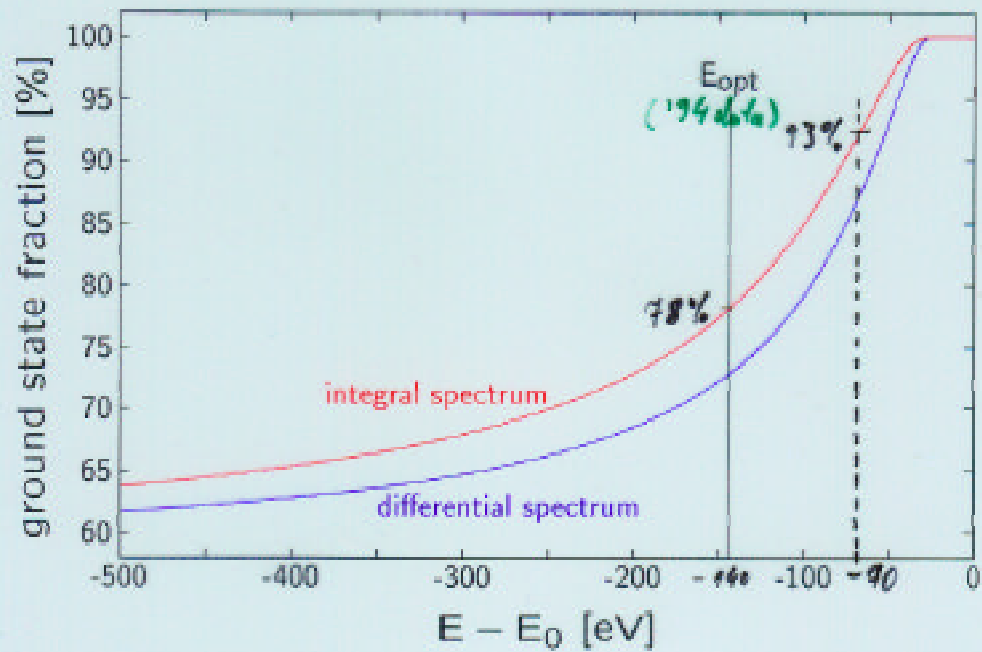
as function of point of measurement E

$$\delta m_\nu^2(E) = \frac{2}{c^4}(E_0 - E)\delta E_0$$

Example (1): Introduce an **external** value of E_0 from some other measurement into the fit which is **too large**, then the fit finds: $m_\nu^2 > 0$ (Ljubimov effect)

Example (2): E_0 is determined from fitting the experimental spectrum, but found **too low**, because the fraction of inelastic events has been underestimated than the fit yields: $m^2 < 0$ (Mainz effect and others)

Fraction of Ground State Transitions in T_2 β -Decay

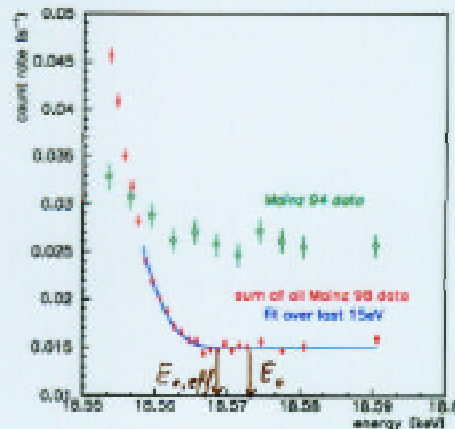


Mainz upper limits on neutrino mass 1

1.) Decorrelate m_ν^2 from systematics:

Sum of all 1998 data (Q3+Q4+Q5) (taken under same conditions)

- Take last 15 eV of β spectrum
 - free of systematics (atomic physics thresholds)
 - no influence from "Troitsk-like" anomaly as observed in Q4
- Add two data points at 18470 eV and 18500 eV
 - to decorrelate m_ν^2 from endpoint E_0 and amplitude
 - minor influence of systematics
 - minor influence from "Troitsk-like" anomalies



$$m_\nu^2 c^4 = -0.1 \pm 3.8 \pm 1.8 \text{ eV}^2 \quad (\chi^2/\text{d.o.f.} \approx 1)$$

$$\rightarrow m_\nu c^2 \leq 2.9 \text{ eV} \quad (95\% \text{ C.L., unified approach})$$

Summary

- Present MAC-E-filters have reached mass limit m_ν well below 3eV
- Old puzzle of "negative" m_ν^2 has been solved: tricky excess of energy losses!
- Residual anomalies show up on a much finer scale, so close to the endpoint that they can hardly be caused by misinterpreting energy losses. They seem to vary in time
- Mainz is shortcircuiting this difficulty by choosing for the evaluation either "clean runs" or a data interval $\leq 15\text{eV}$ below E_0
- Present instruments in Troitzk and Mainz will eventually reach the mass limit of 2eV will probably be able to decide whether residual anomaly is a physics or an instrumental effect
- We need stronger instruments to break the astrophysically relevant 1eV limit and to resolve residual anomalies in the spectrum or search for new ones