



Calorimeter Meeting SSCL

May 1, 1992

Abstract:

Agenda, presentation, and attenders of the GEM Calorimeter Meeting held at the SSC Laboratory on May 1, 1992. Agenda items were: Baseline Status; Forward Activities; Discussion on April 21/22 Cost Review and International Contributions; BaF2 Activities; SSCintCal Activities; LAr Activities; Quartz Fiber Calorimeter; SSCL Test Beam Facilities; and Simulations and Test Beam Data.

A G E N D A

of GEM Calorimeter group meeting on Friday May 1-st at SSCL

9 am H.Gordon - Presentation of base line design status (60')

Forward activities (15') Dave Winn

Discussion on April 21/22 cost review and intl. contributions (30')

Engineering for alternative opt.
BaF2 activities (30') H.Newman Mark Rennich

SSCintCal activities (15') L.Sulak

LAx activities (15') H.Gordon

noon Lunch break (60')

1 pm Philippe Gorodetzky (CERN) - quartz fiber calorimeter (45')

G.Yost - SSCL test beam facilities (20')

Simulations and test beam data (H.Ma, J.Brau ...) (35')

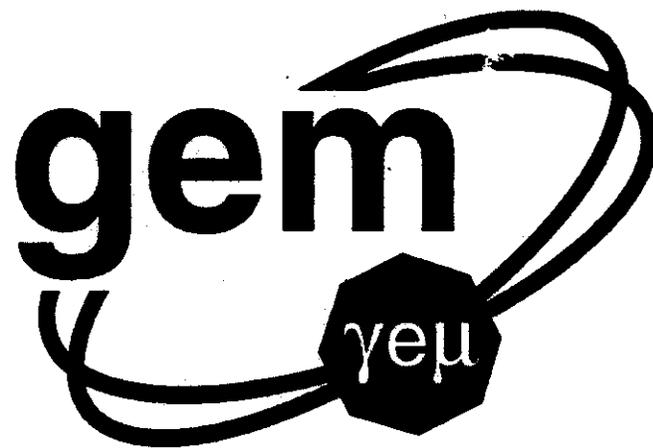
Next meeting, plans, AOB (5')

3:00 pm Adjourn

ORNL>

Bitnet

| | | |
|-----------------------|-----------------|-------------------------------------|
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| KARL VAN BIBBERZ | LLNL | PEGASYS@SLACESA |
| Bennie Ward | Univ. of Tenn | BFLW at SLACVM |



Presentation by:

Yuri Kamyshkov

Cost Review of Calorimetry Options

(SSCL April 21-22)

Preliminary

| | | | |
|------|------------------------------|---------------|--|
| → 1. | BaF2 | \$ 80.3 M | China. \$15 M? |
| → 2. | Sci. Fibers | \$ 55.7 M | Albania \$11 M? Belorus |
| → 3. | Forw. Liq. Sci. | \$ 16.4 M | China ? |
| 4. | Sci. Tiles | \$ 51.8 M | |
| → 5. | LAr e-m + hadr | \$ 136.7 M | (+ 4-5% 91-92 \$) China e-m ? Russia hadr. mo. Korea cryostat |
| 5.1 | LAr separate e-m | \$ ~ 60.0 M | |
| 5.2 | LAr e-m + hadr. forw | \$ ~ 84.0 M | |
| 5.3 | add accordion prerad | \$ > 5.0 M ? | |
| → 6. | Forward LAr | \$ 15.3 M | |
| 7. | LKr parall. plates + π^0 | \$ > 37.8 M ? | (cryostat, install). etc. not included |
| 8. | Si prerad | \$ 16.0 M | |

Comments

- ✓ Baseline LAr (em + hadr + forw) \$ 152 M
 - ✓ Alternative (BaF2 + SciFi + forw) \$ 152 M
 - ✓ LAr hadronic @ R = 360 cm and Z = 550 cm
doesn't satisfy baseline parameters :
only 10.2 λ at $\eta = 0$. and *The problem is
being fixed.*
11.5 λ at $\eta = 3$.
 - ✓ Alternative option @ R = 330 cm and Z = 470 cm
provides 12 λ at $\eta = 0$. and
14 λ at $\eta = 3$.
-
- ✓ SciFi engineering progress was demonstrated
 - ✓ Cost of BaF2 can potentially go up if individual
crtystal treatment will be required
 - ✓ LAr e-m accordion end cap design - no progress

Procedure

Charge from collaboration management to develop the decision process which should include considerations how different cal. options impact the cost of overall detector (via magnet, muon system, assembly etc.)

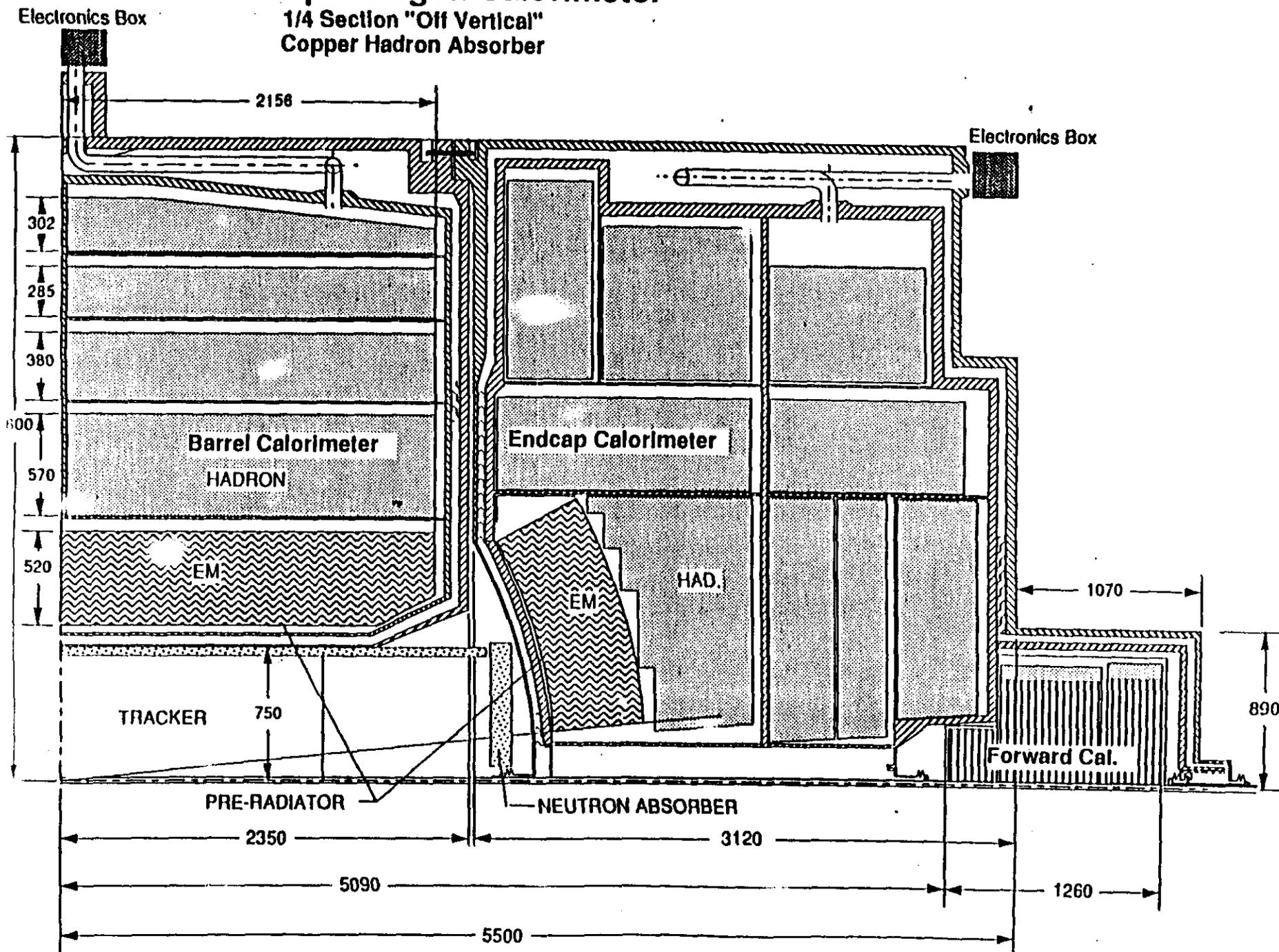
Engineering aspects of saving :

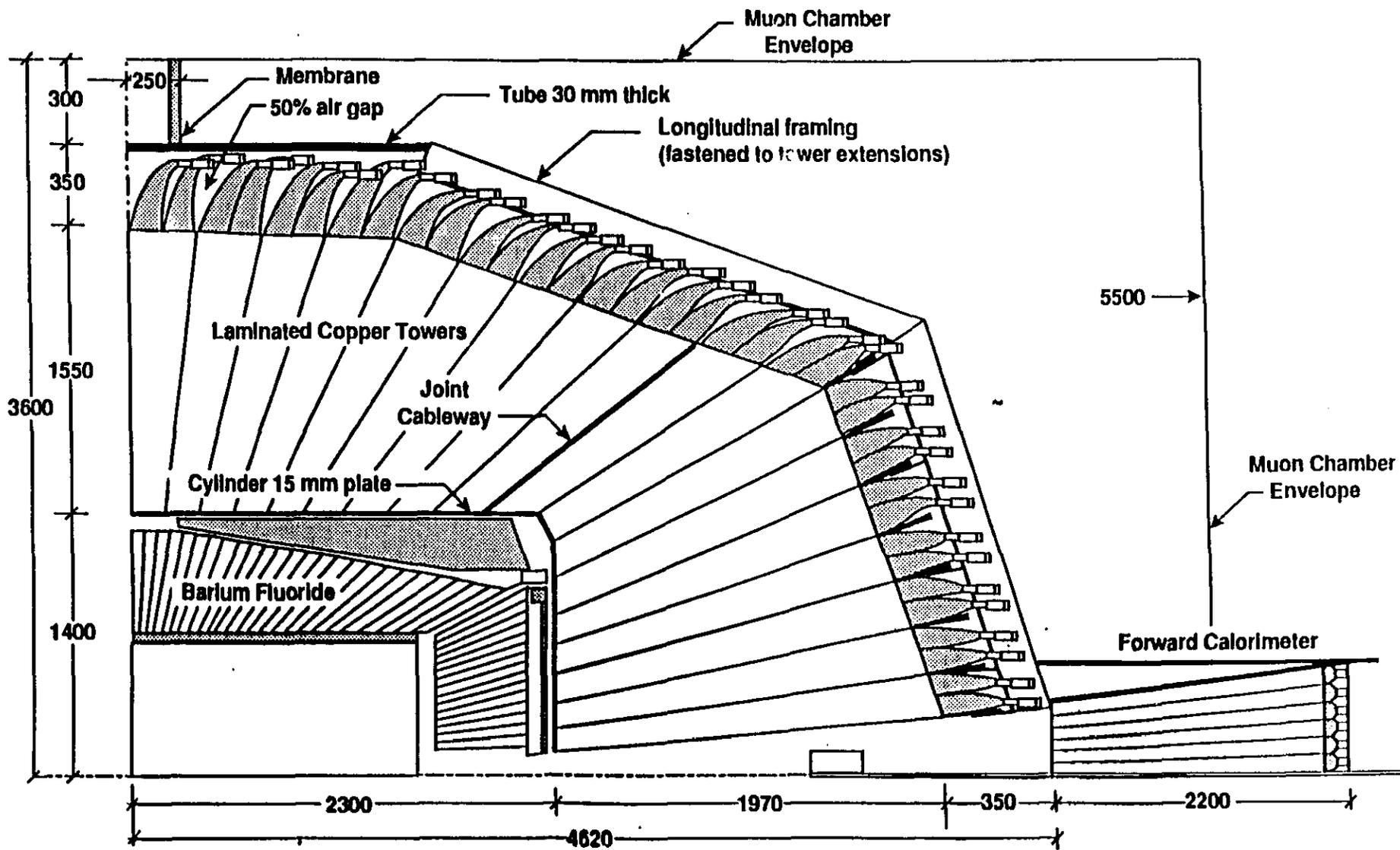
1. Direct cost
2. Detector size
3. Utilities
4. Engineering resources

- ✓ We have set the requirements for two e-m options to be resolved before the decision in August
- ✓ Do we have any new information in hadron calorimetry before August?
- ✓ Are the decisions on e-m and hadron calorimetry are tightly coupled together? Why? *Mixed options.*
(physics, eng., cost, or others?)
- ✓ Do we want NOT to make any decisions between now and August?

GEM Liquid Argon Calorimeter

1/4 Section "Off Vertical"
Copper Hadron Absorber





Dimensions in millimeters

GEM Scintillator Calorimeter System

E-m

- More engineering efforts (+? money) are needed for LAr e-m accordion option
- Engineering aspects of the separate e-m LAr calorimeter option should be studied (M. Rennich, L. Mason)

A. Requirements for BaF2 technology :

1. Demonstrate substantial improvement in radiation resistance of large BaF2 crystals (20-25 cm long) towards the GEM specifications - reach absorbtion length of at least >60 cm at 220 nm after 1MRad irradiation with photons, and if possible high energy hadrons. Present a detailed plan to obtain final GEM quality crystals, along with evidence of manufacturability and cost, including work required to prepare crystals after delivery.

by August 1, 1992

2. As proposed by the expert panel, produce small radiation-hard crystals to demonstrate there are no fundamental limitations in making rad hard BaF2 crystals. (eg. absorbtion length \geq 95 cm at 220 nm after 1 MRad)

by August 1, 1992

3. Address in detail questions of preradiation, wrapping, residual non-uniformity, etc in crystals we can practicably expect to manufacture. Cosmic ray transverse measurements in produced crystals could provide useful data. Provide detailed practical plan for calibration of BaF2 system in-situ :describe calibration strategy, RFQ layout, required calibration time for each proposed technique to achieve necessary accuracy.

by July 1, 1992

4. Show by MC and by lab.tests that the following effects do not dest y the resolution of the BaF2 system (maximum tolerable constant term is 0.6%) :

- residual non-uniformity (as installed);
- non-uniformity developed by possible further radiation damage of "saturated" crystals and/or by possible annealing;
(note - the expert panel and executive committee are not convinced of the proposal to preradiate the crystals)
- accuracy of intercalibration (see point 3);
- short term instabilities of readout system;
- linearity, linearity calibration and dynamic range of readout system.

by July 1,1992

B. Requirements for LAr-Kr technology :

1. Demonstrate by beam tests stochastic term in resolution for non-projective geometry $\leq 7\%/\sqrt{E}$. Determine angular dependence of this resolution.
by August 1, 1992
2. Produce detailed mechanical design/analysis of e-m barrel and end caps with optimization of gap between barrel and end cap, wall thicknesses, etc.
by July 1, 1992
3. Demonstrate by MC simulations for realistic projective geometry and full angular range (between 90 and 5.7 degrees) the resolution
 $\leq 7\%/\sqrt{E} + 0.4\%$
and physics consequences of the gap between barrel and end cap, wall thicknesses, etc.
by July 1, 1992

Hadronic section

- Hadronic LAr cost:

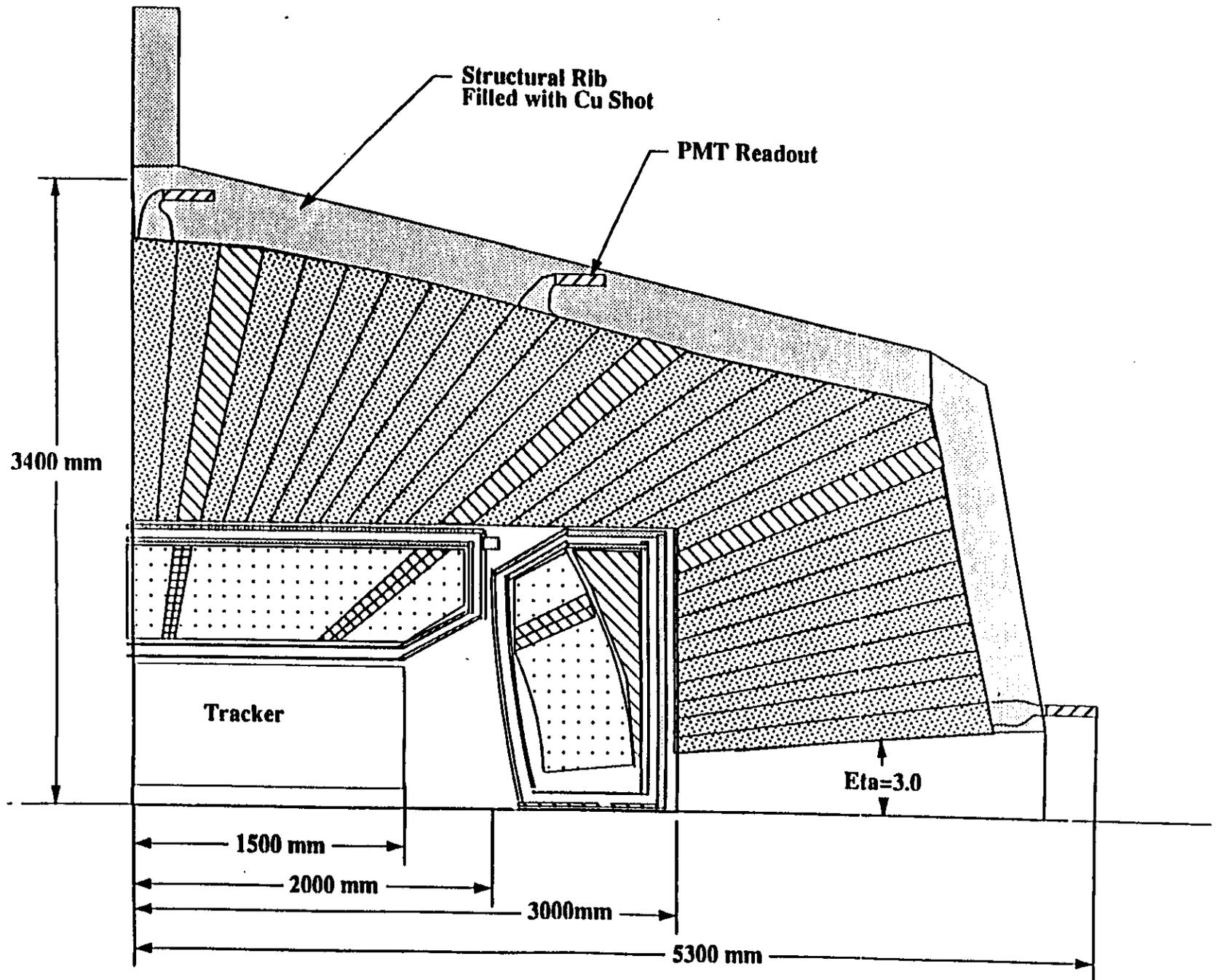
$$\$ (136.7 - 60) \cdot 1.04 \approx 80 \text{ M}$$

If lambda shortage will be fixed

cost most likely will go up

- Hadronic scint. cost:

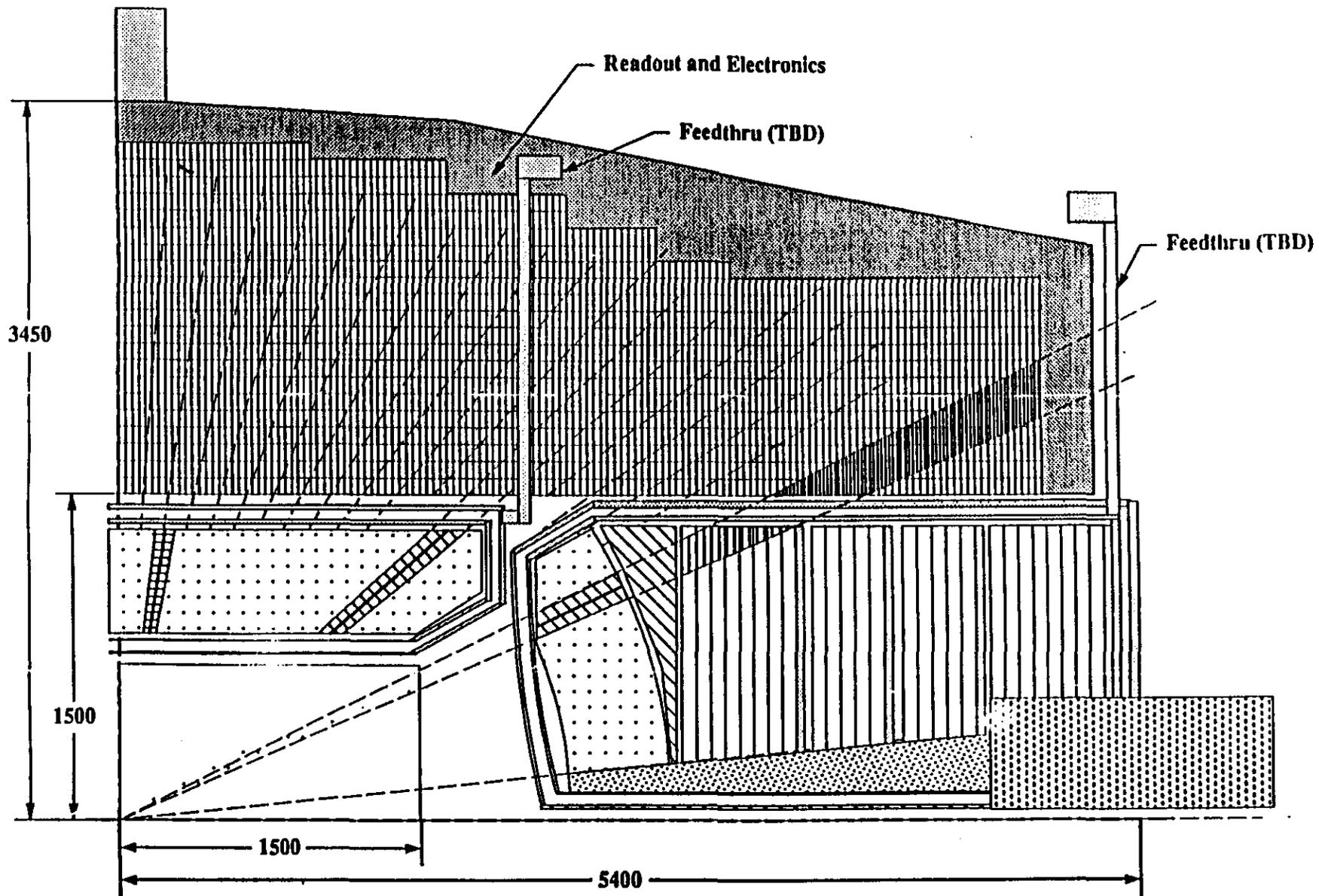
$$\$ 52 - 56 \text{ M}$$



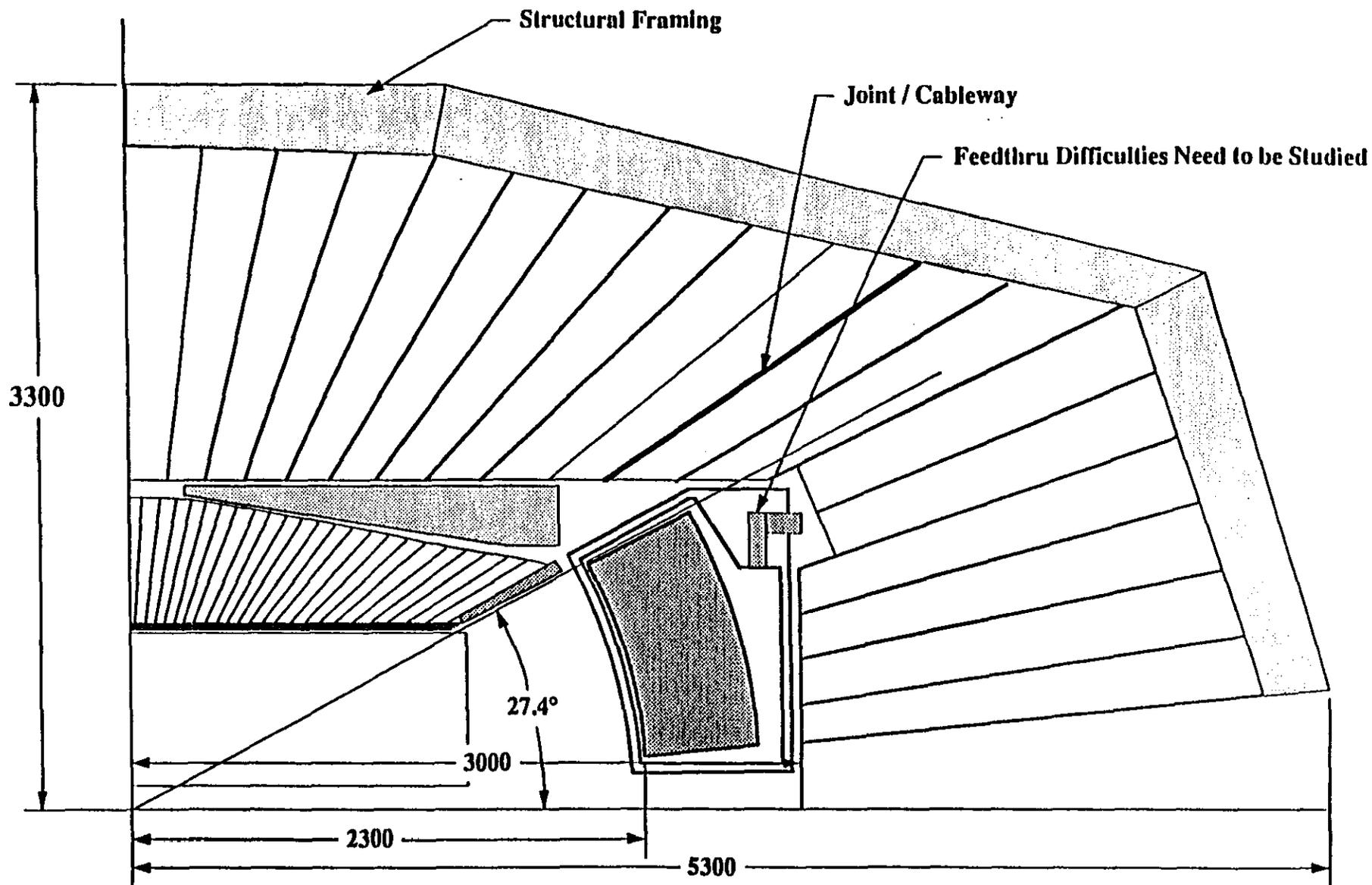
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Spaghetti Hadron Calorimeter
 Copper Fill/0.12 Segmentation
 Liquid Argon EM Calorimeter

Rennich



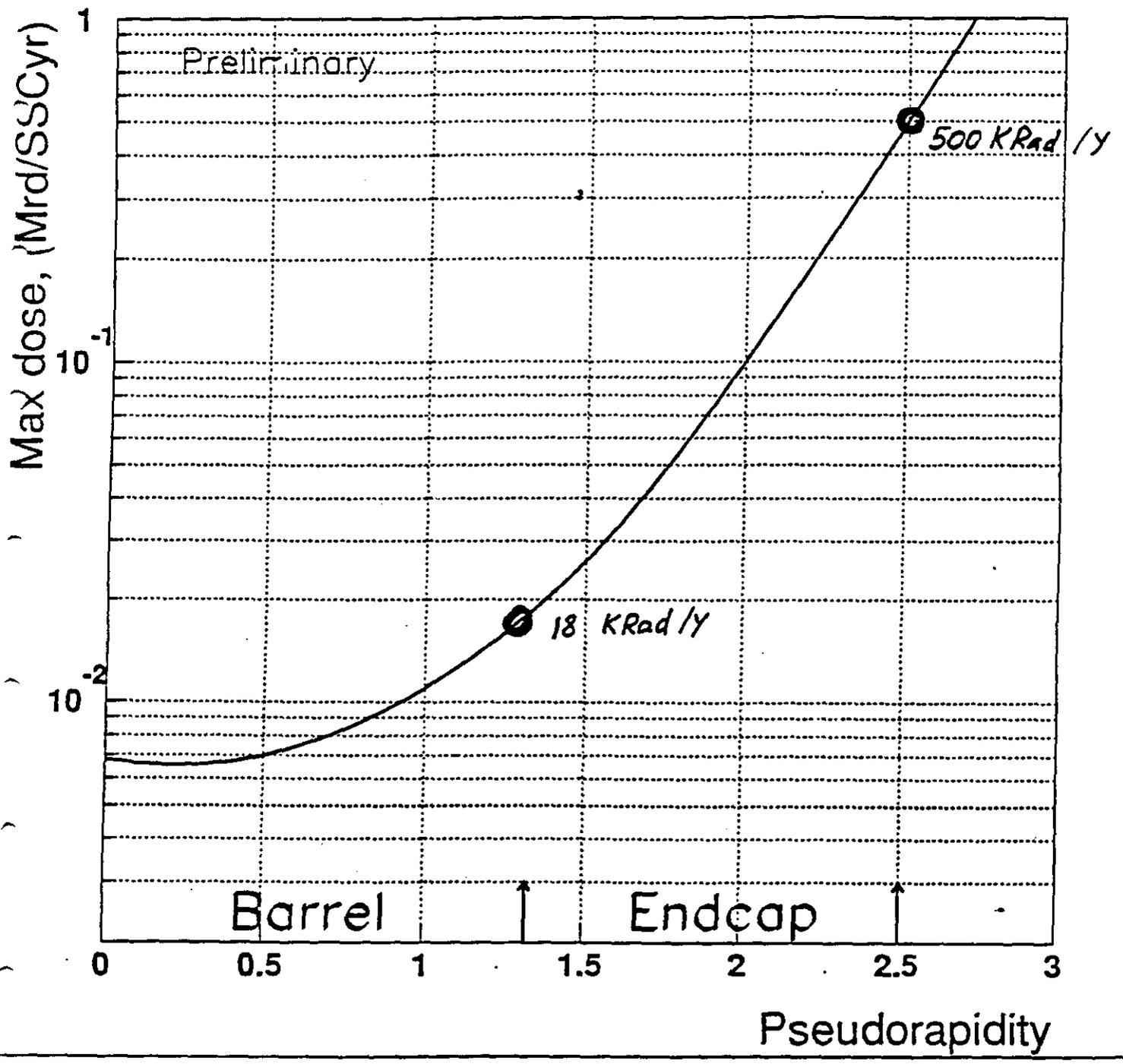
**GEM Detector
 Plastic Tile/Copper
 Hadron Barrel Calorimeter,
 LAr EM - Hadron End Cap and
 LKr EM Barrel Calorimeter**



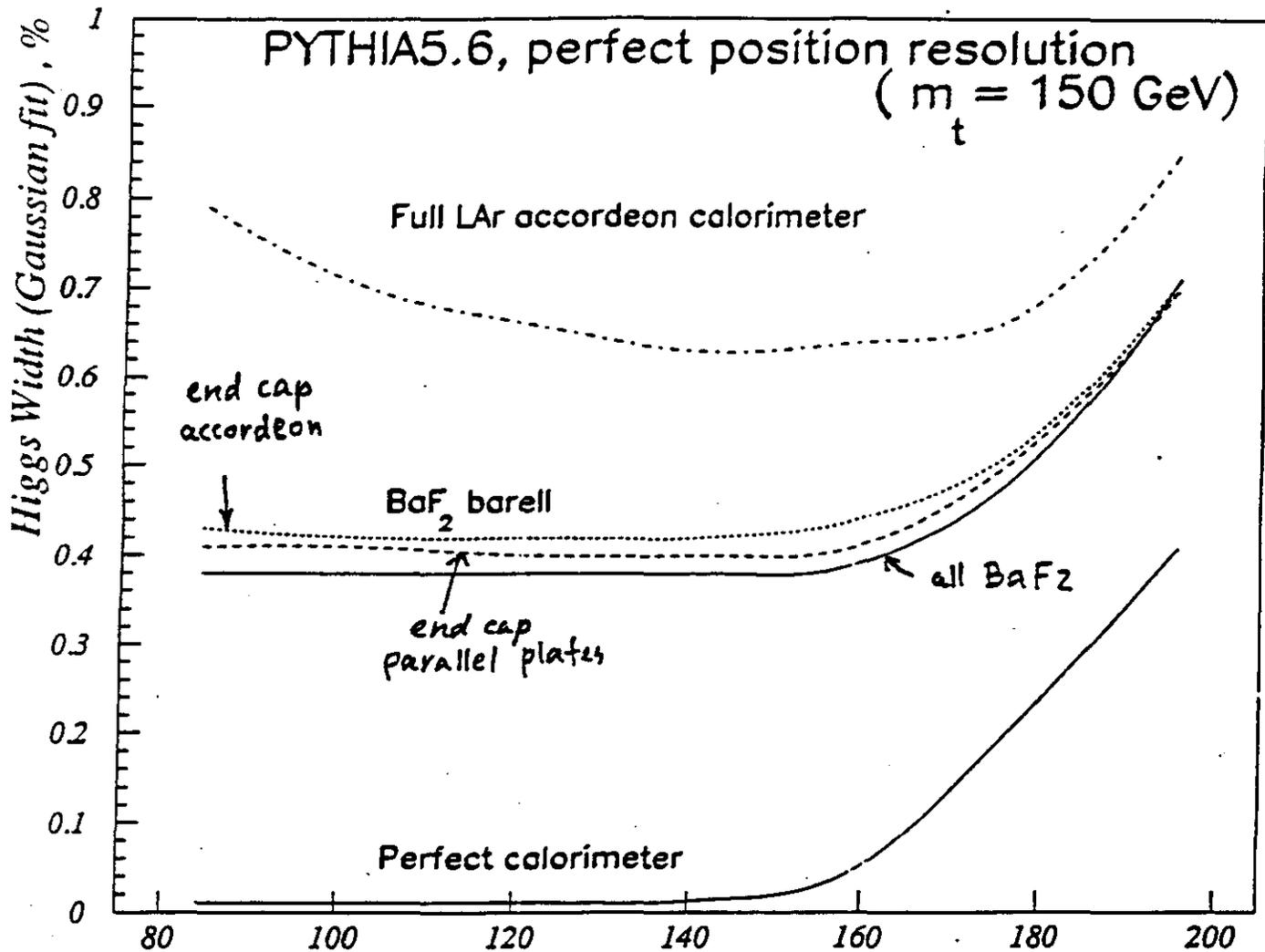
**GEM Detector
BaFFiLAr Calorimeter**

@ 10^{33}

BaF2 DOSE AT CASCADE MAX



HIGGS MASS RECONSTRUCTION



HIGGS MASS (GeV)

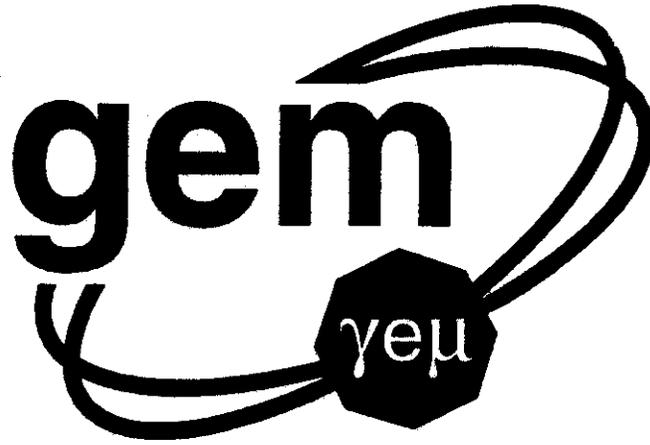
- BaF₂+LAr accordion endcap
- - - - BaF₂+LAr plates endcap
- BaF₂ full calorimeter

Engineering meetings

~ May 12-14 at MMA Denver

May 19 at Boston CDL

~ May 27-28 at SSCL



Presentation by:

Dave Winn

SCCintCAL Forward Calorimetry

STRATEGY:

[1] Fiber Geometry:

No Active Elements, No HV in high radiation environment

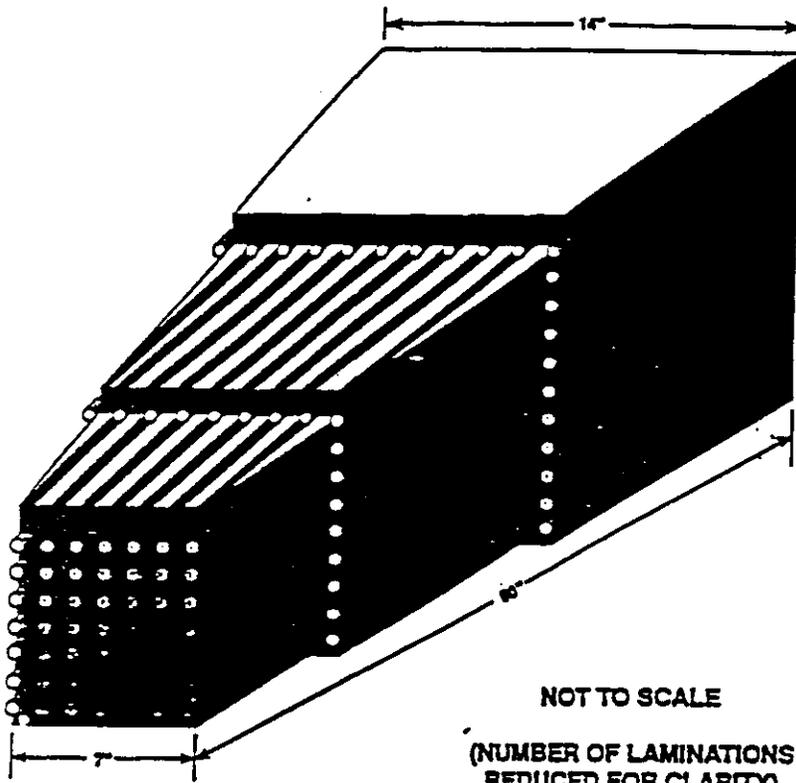
Fast - Scintillator, Cerenkov+no WLS+optics -> 40 ns Eh, 15 ns em

PMT survives > 100's MRad, also shielded by calorimeter

Replacement possible: fibers exit back of calorimeter

TYPICAL COPPER LAMINATED TUBE AND PLATE TOWER

AS ASSEMBLED, BEFORE FINAL MACHINING



SIDE VIEW

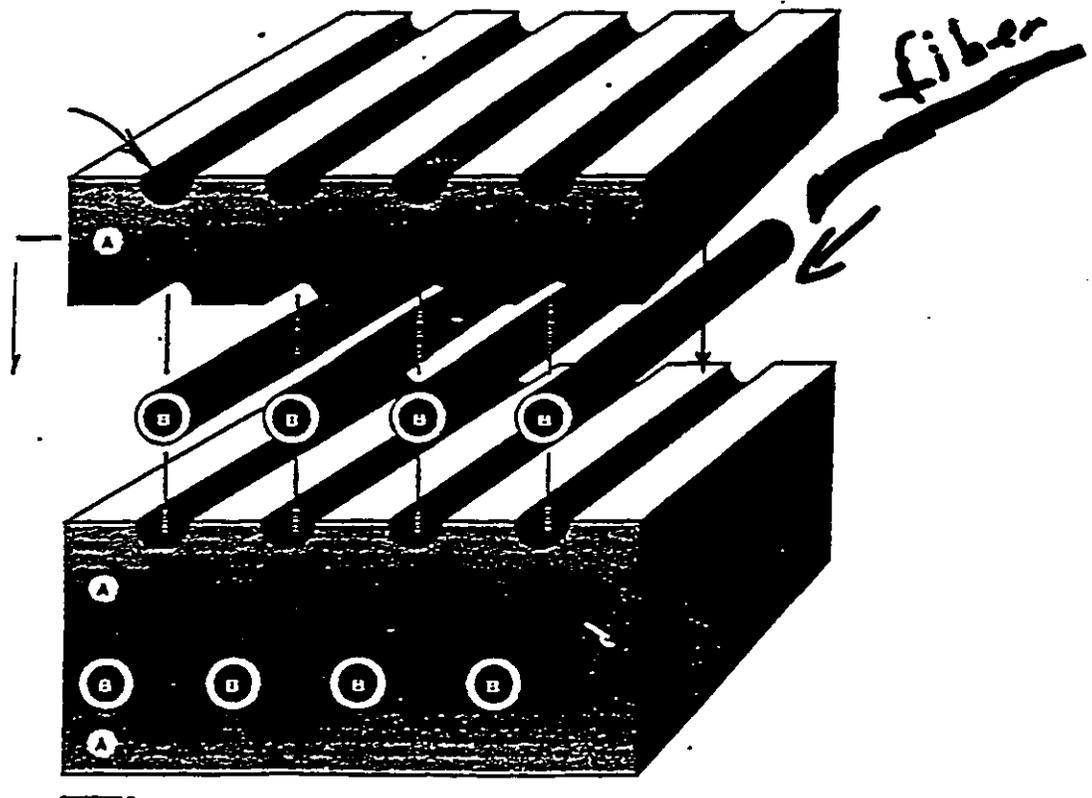
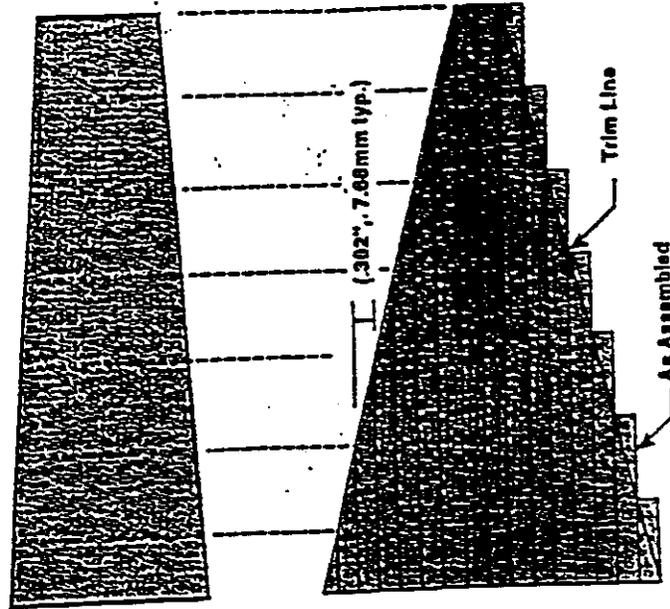
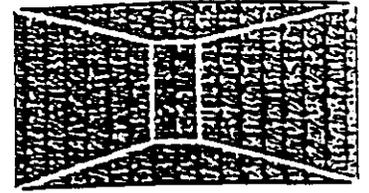
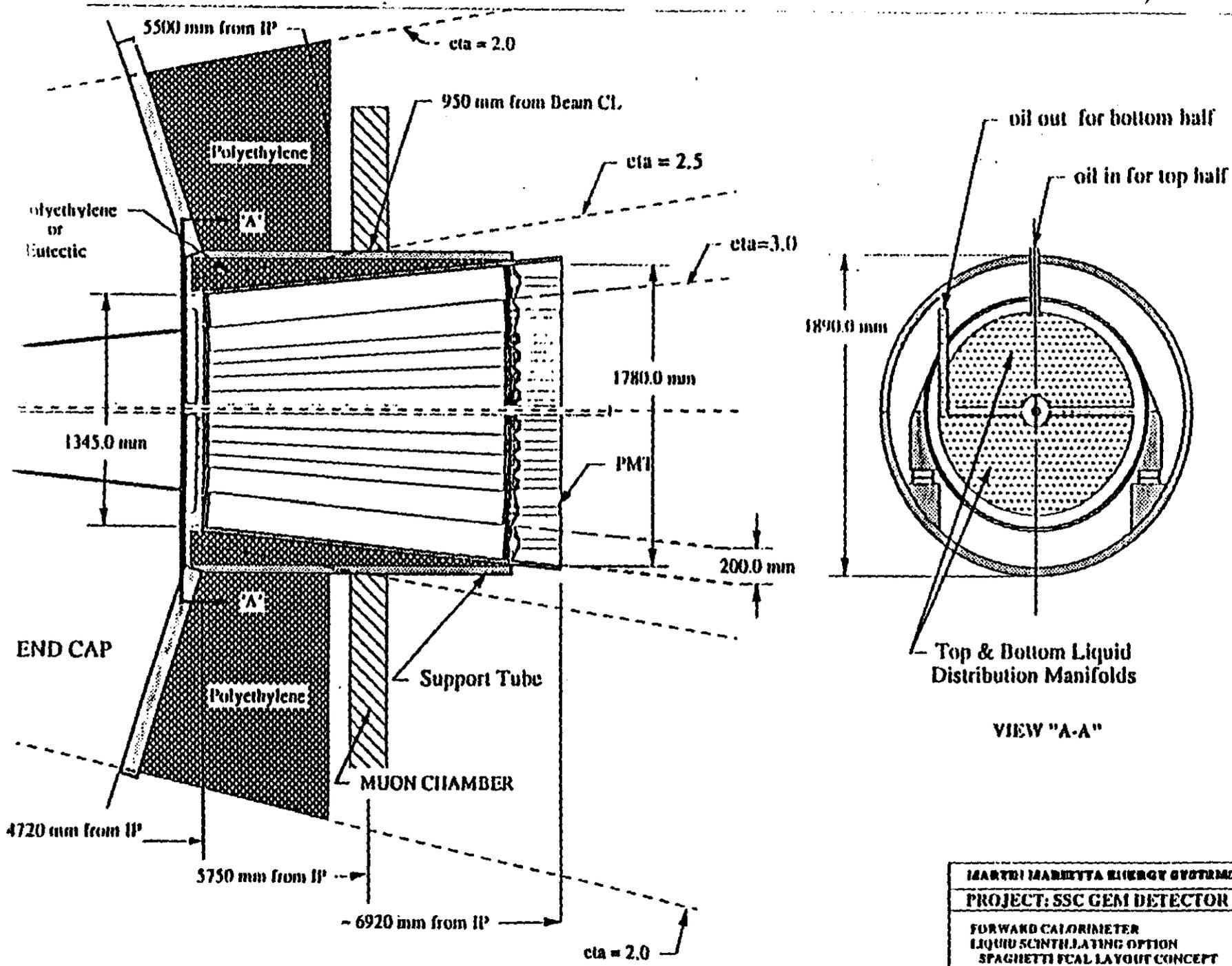


Figure 15a,b: Schematics of SSCintCAL Cu hadron towers - example only for use of the fibers.



NOTE:
1. THE PHYSICAL SYSTEM IS NOT SHOWN.

| | |
|---|---------|
| MARTIN MARIETTA ENERGY SYSTEMS | |
| PROJECT: SSC GEM DETECTOR | |
| FORWARD CALORIMETER LIQUID SCINTILLATING OPTION SPAGHETTI FCAL LAYOUT CONCEPT | |
| PRINT NO: D-FCLS-MD001 | REV 3 |
| DRAWN BY: S. M. CHAE | 3/31/92 |
| CHECKED & APPROVED BY: | |

[2] Two General Rad-Hard Alternatives to Plastic Fibers

•Liquid Core Scintillator Fibers

Liquids:

- Existing liquid scintillators rad-hard to >200 MRad
- pumpable/replacable
- Isopropyl biphenyl: reactor coolant, no plating, $n=1.58$
- methyl naphthalene, $n=1.62$; benzyl alcohol, $n=1.55$

Claddings:

- radiation damage -> Labs, not to refractive index
 - PEEK ~ 0.5-1 GRad, $n\sim 1.5$
 - Kapton ~ 0.5-1 GRad, $n\sim 1.55$
 - Halar ~ 200 Mrad, $n\sim 1.43$
 - PUR ~ 200 Mrad, $n\sim 1.5$
 - Quartz or glass tubing, $n\sim 1.5$
 - Viton, Polysiloxanes, $n\sim 1.4-1.5$

Commercial Technology - Oriel Inc.

- N.A.=0.47, Labs=5m, 3 mm fiber polymer spaghetti
- Sealed optical end plugs
- Flexible (can be bundled)
- Rugged
- Replace Claddings & Liquids as above
- Optical surface obtained by standard die extrusions

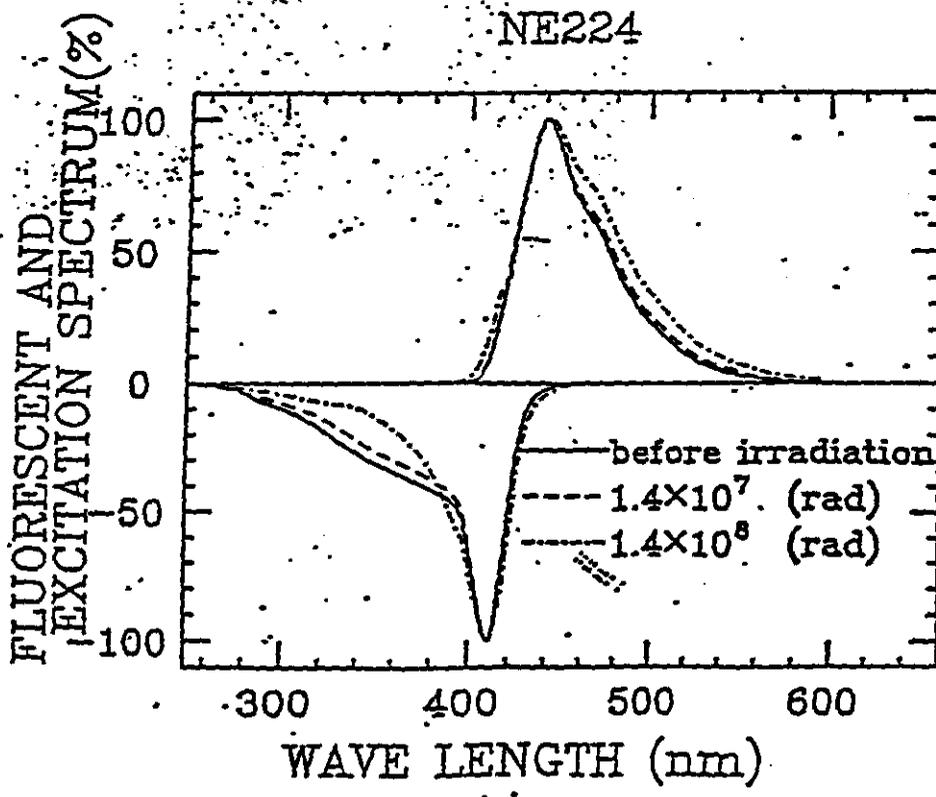
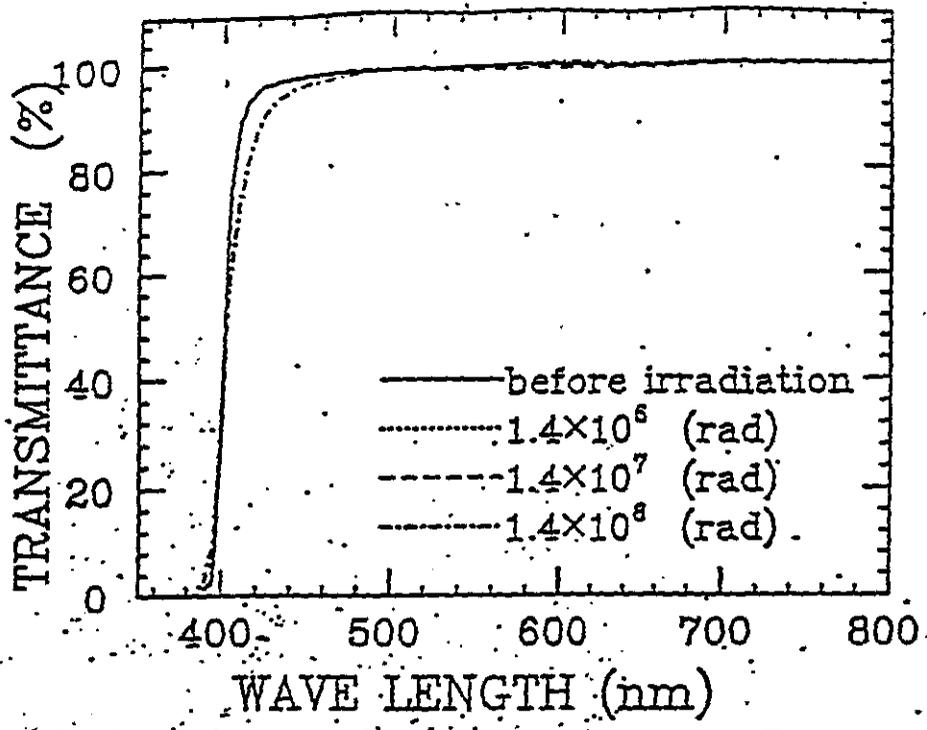


Figure 1: Typical toluene based liquid scintillator's (NE224) properties before and after 140 MRad [Chiba et al p 381, CDF Conference at Tsukuba, April, 1990, KEK publ.]

HALAR® ECTFE FLUOROPOLYMER

HALAR® ECTFE is a melt processable fluoropolymer from Ausimont, Inc. It possesses a unique combination of properties as a result of its chemical structure - a 1:1 alternating copolymer of ethylene and chlorotrifluoroethylene.

HALAR fluoropolymer offers excellent chemical resistance, good electrical properties, broad use temperature range - from cryogenic to 300°F (150°C), and meets the requirements of the UL-94 V-0 vertical flame test in thicknesses as low as 7 mils. It is a tough material with excellent impact strength over its broad use temperature range. HALAR ECTFE also maintains useful properties on exposure to cobalt 60 radiation at dosages of 200 megarads. It is one of the best fluoropolymers for abrasion resistance.

HALAR fluoropolymer is a thermoplastic which can be processed by virtually any technique applicable with polyethylene. It can be extruded, injection molded, blow molded, rotomolded, and applied by ordinary fluidized bed or electrostatic coating techniques. It is available in a range of viscosity grades for extrusion and molding applications. HALAR ECTFE powders are available in three different particle sizes optimized for specific coating processes.

For The Designer and Processor

Processability - HALAR fluoropolymer can easily be processed by injection molding, compression molding, blow molding, extrusion, rotomolding, and electrostatic spray and fluidized bed coating techniques.

Thermal Properties - The continuous thermal rating for HALAR fluoropolymer is 300°F (150°C) depending on resin grade.

Chemical Resistance - HALAR fluoropolymer is resistant to a wide variety of corrosive chemical and organic solvents, including strong acids, chlorine and aqueous caustic. No known solvent dissolves or stress cracks the polymer at temperatures up to 250°F (120°C).

Electrical Properties - The dielectric constant of HALAR fluoropolymer is low (2.5 for solid insulation to 1.5 for foamed insulation) and stable across a broad temperature and frequency range. The dielectric strength is 2000 volts/mil in 1 mil thickness.

Toughness and Strength - HALAR fluoropolymer possesses excellent mechanical properties over a wide range of temperature from cryogenic temperatures to 300°F (150°C). It has nylon-like durability and provides excellent impact resistance at ambient and subambient temperatures.

Weldability - HALAR fluoropolymer is weldable with ordinary thermoplastic welding equipment at 480-500°F (249-260°C).

Barrier Properties - Barrier properties of HALAR fluoropolymer are excellent - 10 to 100 times better than those of PTFE or FEP to oxygen, carbon dioxide, chlorine gas or hydrochloric acid.

Radiation Resistance - HALAR fluoropolymer maintains useful properties on exposure to cobalt 60 radiation of 200 megarads.

Flammability - HALAR fluoropolymer, in thicknesses as low as 7 mil, has received a UL 94 V-0 rating. The oxygen index (ASTM-O-2853) of HALAR fluoropolymer is 60 on a 1/16" thick specimen and 48 for .0005 inch diameter filament yarn.

Flame Spread and Smoke Generation - HALAR fluoropolymer, because of its low flame spread and low smoke generation characteristics, has received UL 910 (NFPA 262) listing up to 200 pair communication plenum cable.

Weatherability - No change after 3000+ hours exposure in a Xenon arc weather-ometer.

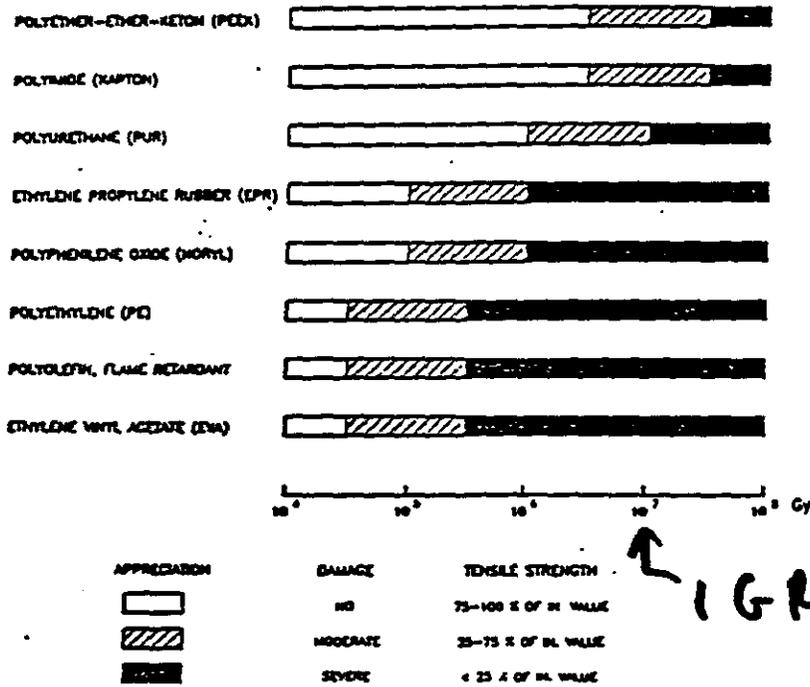
Machinability - Similar to nylon 6.

Water Absorption - Less than 0.1%.

$n = 1.42$

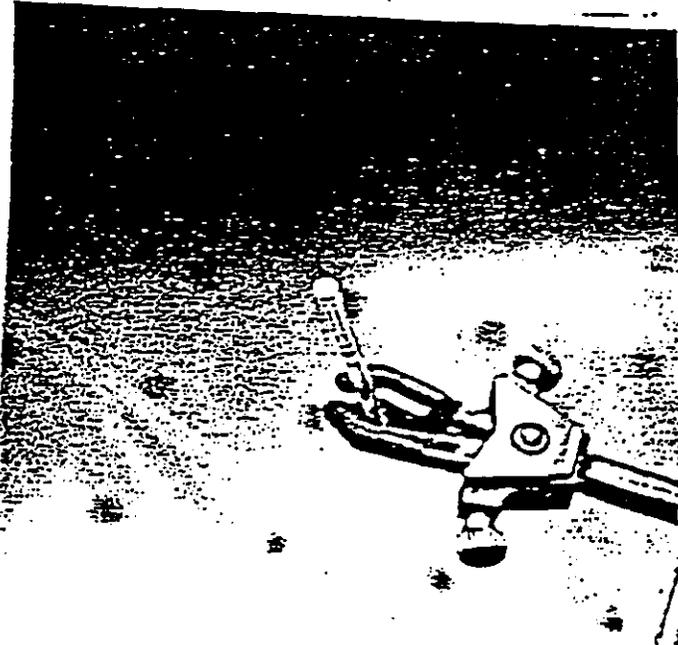
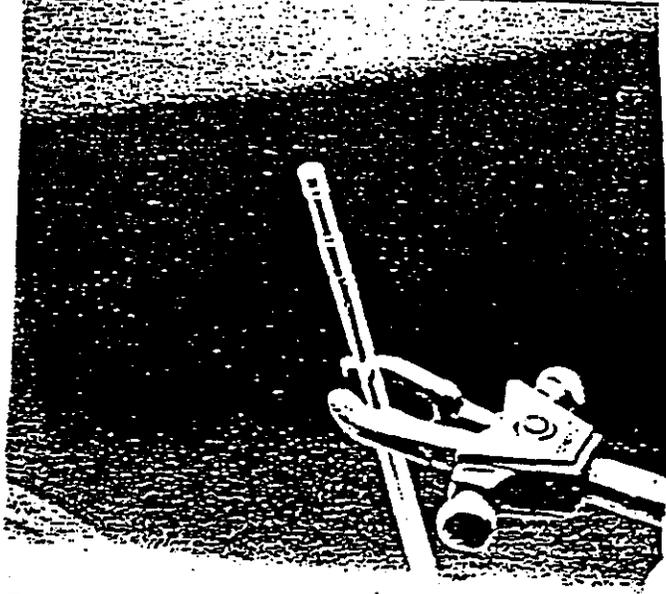
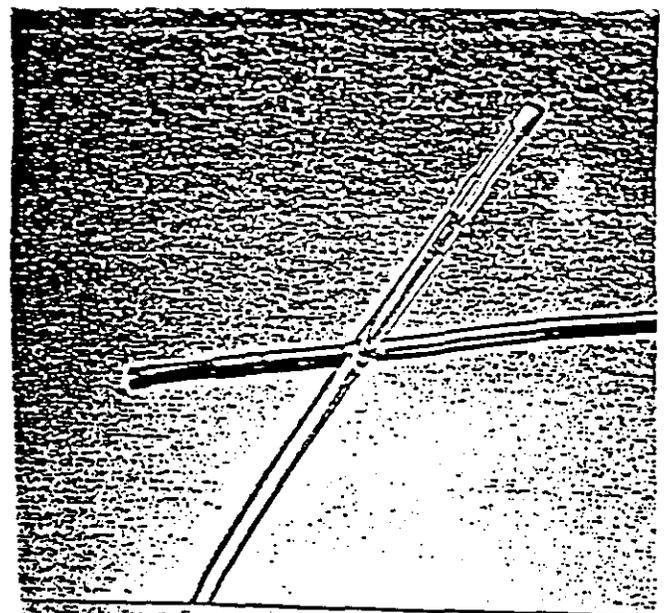
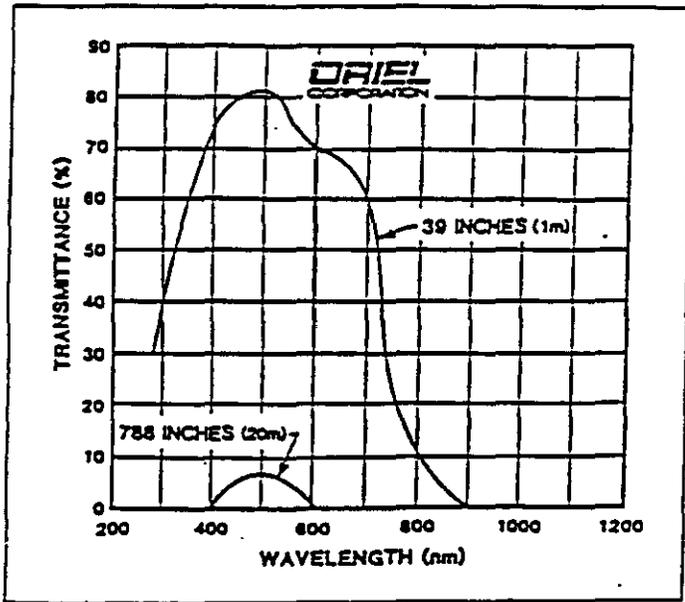
Figure 6: Properties of Halar from the manufacturer; Note Radiation Resistance of 200 MRads.

GENERAL RELATIVE RADIATION EFFECTS
CABLE INSULATION AND SHEATH MATERIALS



ENVIRONMENT : AIR + AMBIENT TEMPERATURE

Figure 7: Radiation resistance of some polymers from a CERN report. Note the highest 3 will be investigated as claddings using commercial tubing manufacturers as the tubing sources.



Oriel LIQUID LIGHT GUIDES

- Excellent light gathering - large cores, no packing losses, high N.A.
- High UV transmittance - from 270 nm
- Flexible tubing

Liquid Light Guides transfer ultraviolet and visible radiation very efficiently. These light guides function in a similar manner to fiber optic bundles, using internal reflection to guide energy along their length, but they have higher transmittance than fused silica bundles of equivalent core size and are more flexible. (See Fig. 1 for a transmittance curve.) Liquid light guides are usable from 270 to 720 nm.

CONSTRUCTION

Liquid Light Guides consist of a plastic tube covered by a protective aluminum spiral and covered by a PVC jacket. The inner tube is filled with a proprietary, transparent, anaerobic non-toxic fluid. The tube is sealed at both ends with polished fused silica windows and protected by an interlocking stainless steel sheathing.

Like our fiber bundles, these guides have 11 mm female adapters that mount to any Oriel Fiber Optic Cable Accessory found on pages 8-20 to 8-23.

Our liquid light guides are robust and long lived in normal use. To ensure long life the temperature at the input face should not exceed 60 °C.

LARGE CORE DIAMETERS

We offer liquid light guides in larger core diameters than our fiber bundles. The guides use the full input aperture and have no packing fraction losses. The large core diameters, and the high numerical aperture allow efficient coupling to large light sources. Core diameters of 3 and 5 mm are standard; 8 mm core diameters are available on special order.

LIGHT ACCEPTANCE

The acceptance angle α for these guides is 28 degrees, with corresponding NA of 0.47. The acceptance cone angle 2α is 56 degrees. The geometrical extent, a measure of fiber optic light gathering capability, is $14.4 \text{ mm}^2 \text{ sr}$ for the 5 mm guides. This compares with 3 for the 3.2 mm diameter fused silica bundles and 7 for the 3.2 mm diameter glass bundles.

BEND RADIUS

The minimum bend radius for the 3 mm core guides is 3 inches (76.2 mm), and 4.8 inches (122 mm) for the 5 mm core guides.

Changing the position of the guide will change the output beam distribution and power. Care should be taken to avoid appreciable movement of the guide during a measurement.

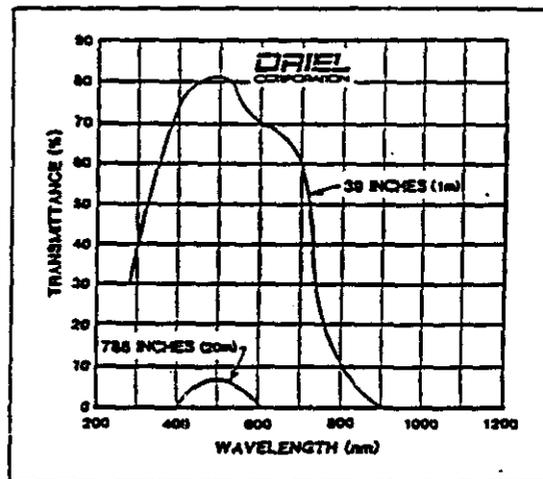
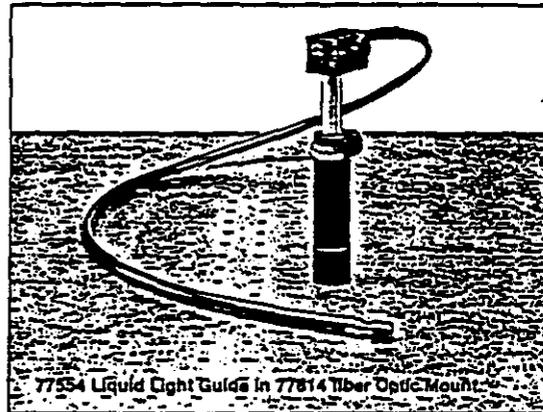


Fig. 1 Transmittance of Oriel Liquid Light Guides.

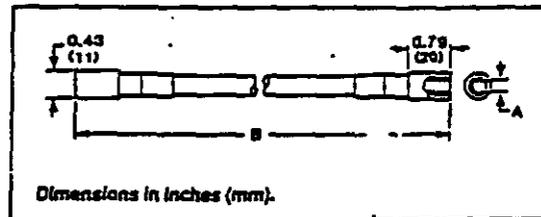


Fig. 2 Oriel Liquid Light Guides.

Figure 8: Oriel spec sheet for liquid optical light guide. Note sample in photograph is armored.

ORIEL

CORPORATION

April 1, 1992

Dr. David Winn
Fairfield University
Physics Department
Fairfield, Connecticut 06430

via fax: 254-4126

Dear Dr. Winn:

We are looking forward to our participation with your team in the development effort for the SSC forward calorimeter based on liquid light guide scintillator concept.

The time scales listed in your proposal, even though somewhat ambitious, appear to be feasible and we would exert every effort to help you meet them. The total budget for our part of the effort, with some extra contributions of our engineering team time, appear to be appropriate for the scope of the project. We would like to offer to contribute extra effort as needed, up to 20% of that planned below, to help achieve the goals of this work.

The project will be charged at cost for materials, components, and consultant time. Engineering and assembly time will be charged with 35% corporate overhead rate added. We are projecting that materials and fixtures will consume \$29,200. Use of the equipment located in Oriel's test laboratories, such as the light sources and detector systems, is included in the overhead rate. We plan to utilize the consulting services of Mr. Clyde Hinman for 120 hrs, at an hourly rate of \$40.00, for a total expenditure of \$4,800. A total of 400 hrs of labor is planned for the assembly and testing of the 800 fibers for the Calorimeter Tower. 600 hour effort is planned for adapting our light guide technology to this new application and assistance in materials testing and optical properties determination.

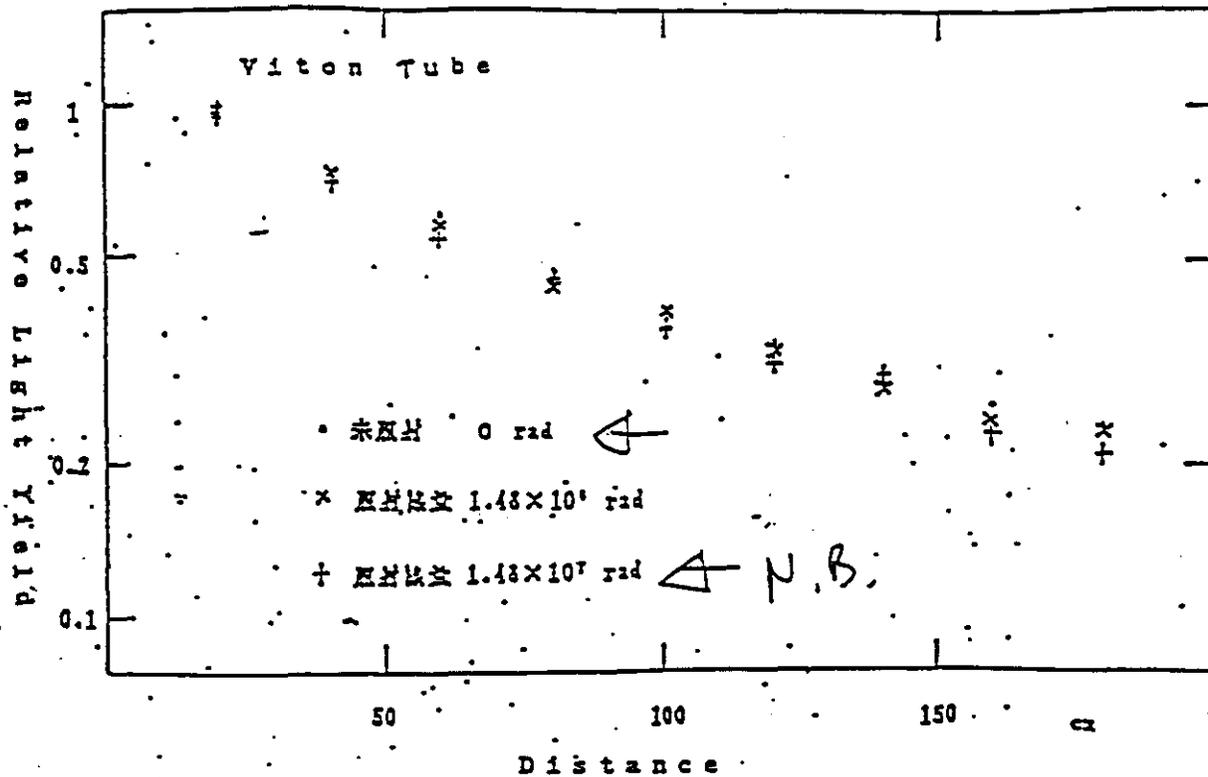


Figure 3a: Light yield as a function of distance, before and after 15 MRads: NE224 in a Viton Tube [Chiba et al p 381, CDF Conference at Tsukuba, April, 1990, KEK publ.]

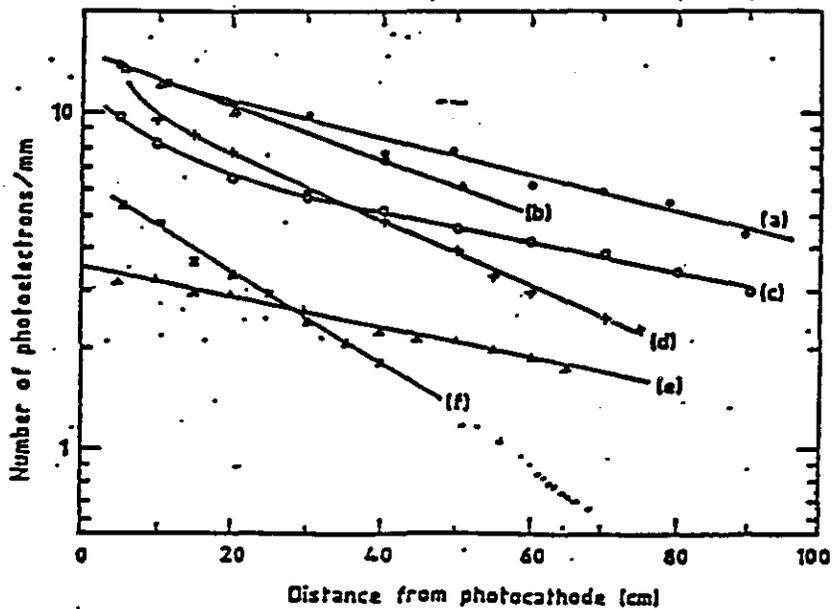
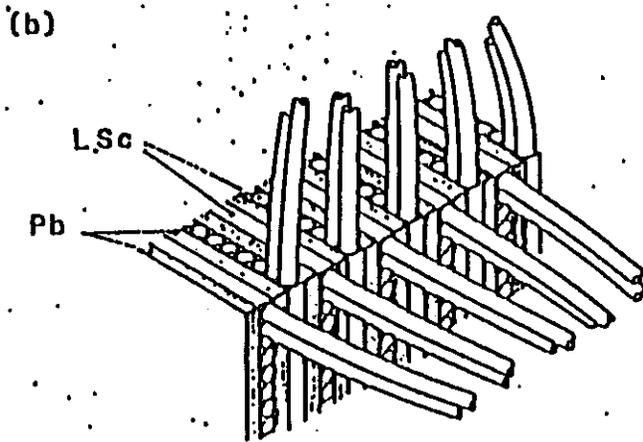
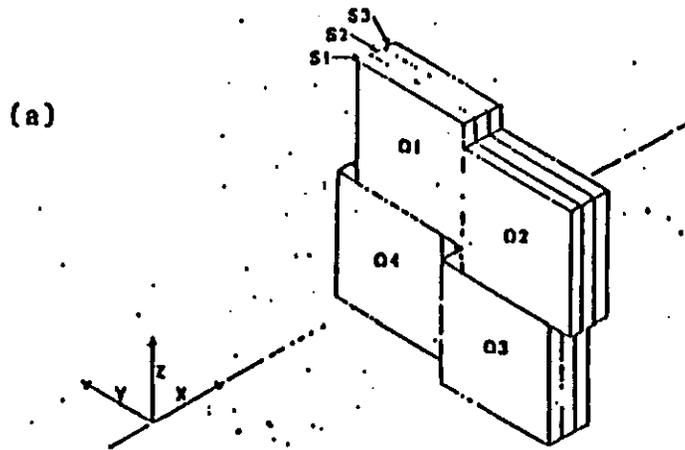


Figure 3b: Light yield as a function of distance in glass capillaries for various high index liquids produced by Bicron, Inc. [J. Bahr, et al., NIM A306 169 (1991)]



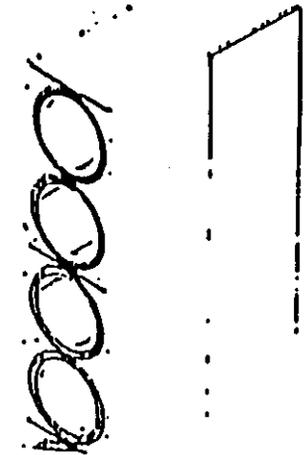
The geometry of the calorimeter, showing the four quadrants (Q) and the three segments (S). (b) The structure of the outer corner of one segment showing the lead (Pb) and the tubes of scintillator (Sc). Light from groups of tubes in the five layers in a given direction was mixed and fed to one photomultiplier. Groups of two tubes gave a granularity of 1.07 cm.

Fibreglass bonds

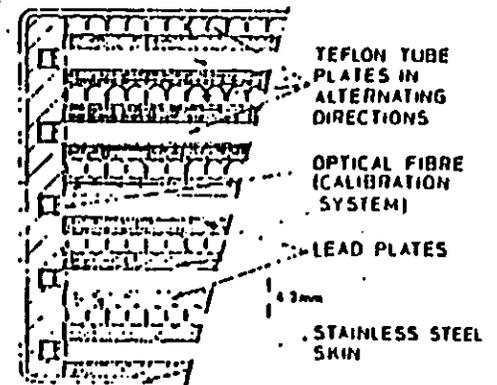
Stainless steel bonds

Teflon tube

Epoxy



Teflon tube plate.



Calorimeter construction: detail of one segment.

Figure 5: Plate calorimeter using plates of liquid scintillator in teflon tubes, used at CERN [M. Bonesini et al NIM A270 32 (1988) & NIM A263 325 (1988)]

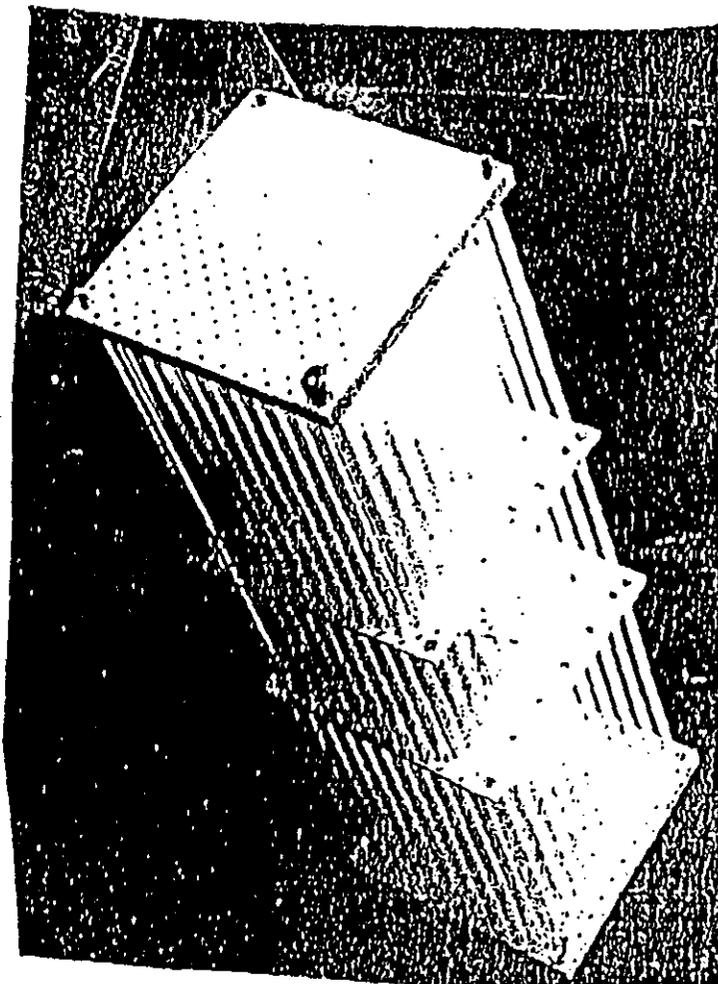
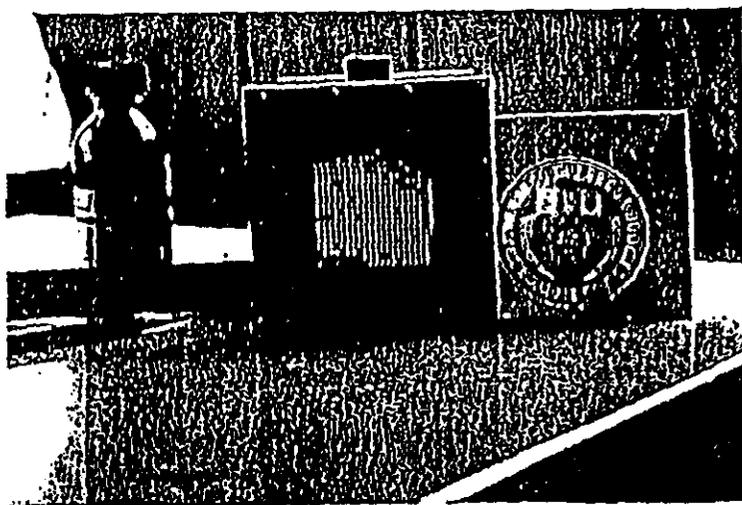
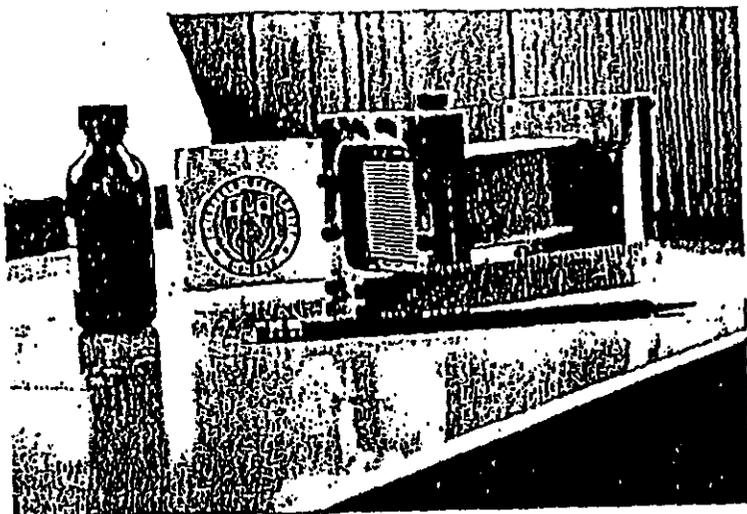
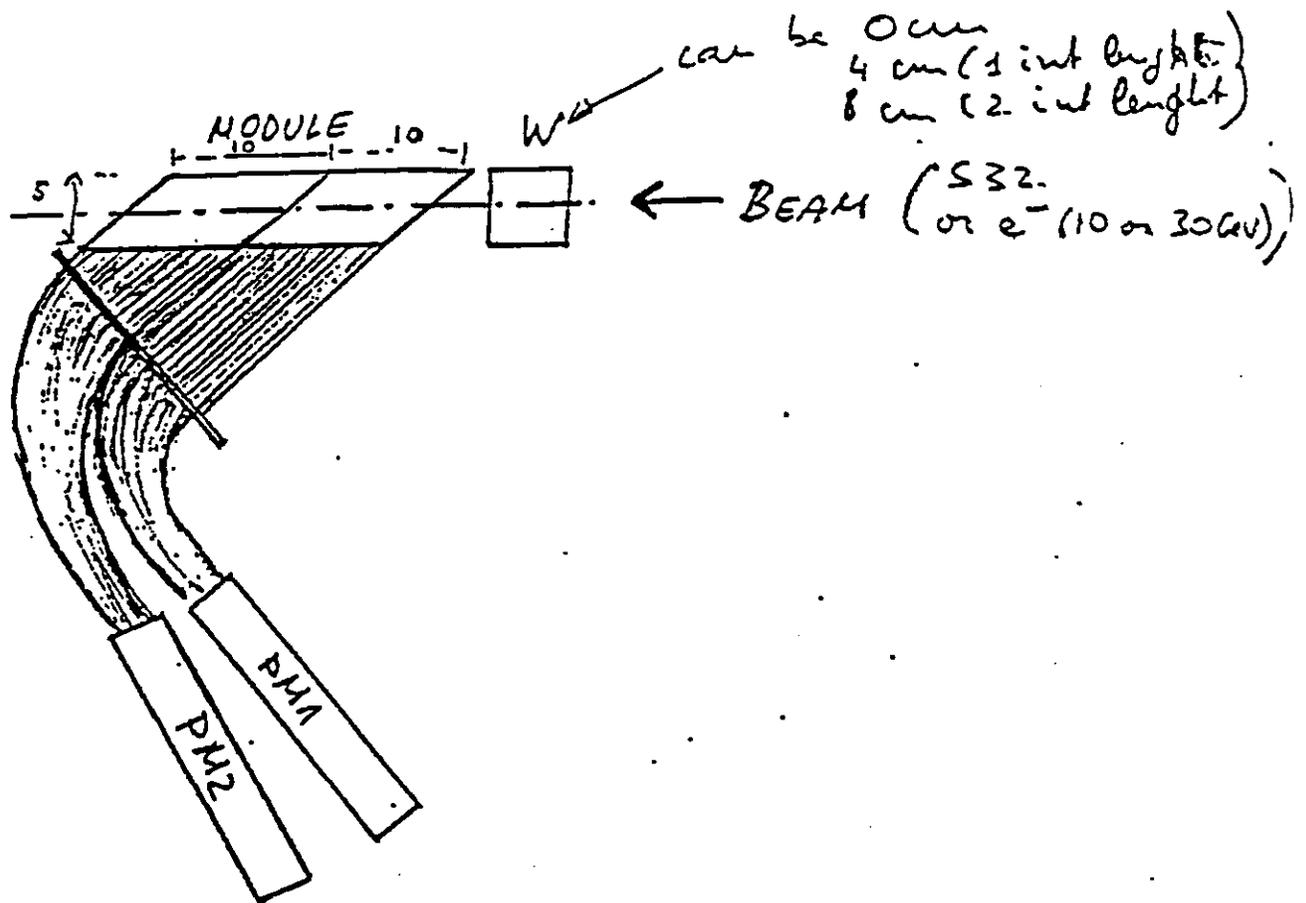


Figure 4: Liquid spaghetti E-M calorimeter using teflon tubes filled with liquid scintillator, constructed and tested at BNL by the authors [IEEE TNS]. (a): Teflon spaghetti armature; (b) Cast Pb block encased in SS containment vessel with glass windows for PMT. The resolution was about $17\%/E^{*0.5}$ at low energy (1-3 GeV).

- Quartz Fiber Cerenkov Sampling

- [Developed for NA38 - Ph. Gorodetzky]

- Potentially hard to 10's GRads
 - 1 mm Quartz fibers w/ PEEK cladding: \$5/m
 - 3 mm Quartz tubes: \$9/m
 - Energy: 3-4 p.e./GeV hadrons; 35-40%/√E em
 - Resolution: ~8-10% constant term hadrons
 - Exceptional Speed



DIMENSIONS OF THE MODULE: $5 \times 5 \times 20$ cm

THICKNESS OF TUNGSTEN BLOCK: 0, 4, 8 cm

1474 FIBERS 50 cm LONG.

DISTANCE BETWEEN FIBERS 2.22, $V_{PB} : V_{FIB.} = 4:1$

INTERACTION LENGTHS FOR S^{32} :

- IN W $\lambda_{int} = 4$ cm

- IN MODULE $\lambda_{int} = 9.1$ cm (including quartz fibers)

Figure 10: Schematic of the the $5 \times 5 \times 20$ cm quartz fiber Cerenkov sampling calorimeter module read out by 2 PMT, tested at CERN in an Sulphur-32 (160 GeV/nucleon) or e^- (10-30 GeV) beam. Note that the 1mm quartz fibers (20% p.fraction) are arranged to maximize at least some of the Cerenkov emission along the fiber meridional ray. The tungsten is to partially develop the hadron shower before the calorimeter. [Courtesy of P. Gorodetzky].

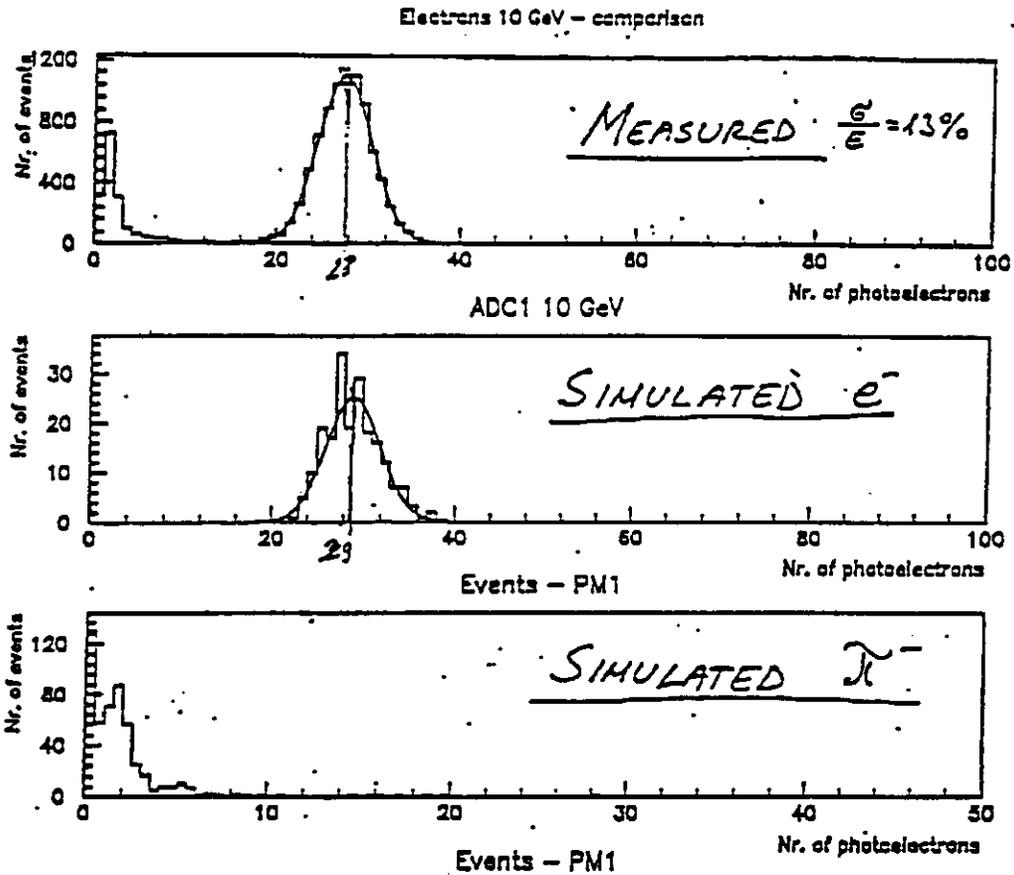


Figure 11: The measured & simulated ADC spectra (resolution) for electrons at 10 GeV in the quartz fiber Cerenkov calorimeter of Fig. 10. Note the simulated 10 GeV pions reproduce the low energy peak in the data. The resolution is roughly $37\%/\sqrt{E}$. [Courtesy of P. Gorodetzky].

PMT Specifications

DRW - 5/1/92

Photocathode:

Photocurrent: 2% linearity for 10^7 p.e. in 10 ns

Lifetime: 1 mC for 50% gain loss

Uniformity: $\pm 10\%$ over 25 mm circle

Multiplier:

Gain: ~6,000-10,000

Pulsed Current: 25 mA in 10 ns with $< 0.5\%$ non-linearity

200 mA in 10 ns $< 2\%$ non-linearity

Risetime (Δf^n pulse, 20%-80%, 10ma in 50 Ω): < 4 ns

Falltime (Δf^n pulse, 80%-20%, 10ma in 50 Ω): < 6 ns

Width at Baseline (5%): < 12 ns

Pulse Shape Change: $< 20\%$ for pulses between 0.1 -100 mA

Average Anode Current: 1 μ A

Lifetime: 2 C w/ 50% loss

Anode Capacitance: < 10 pF to all other electrodes combined

Anode Inductance: < 45 nH

Noise:

Single p.e.: < 100 kHz at 0.5 pe peak

Afterpulsing: $< 0.1\%$ /p.e for afterpulses > 10 p.e.

Pre Pulsing: Arrival < 5 ns early

amplitude $< 0.5\%$ of signal

Stability:

Long Term: $< 1\%$ /month

Hysteresis: $< 1\%$ gain shift within 100 ns after pulse of 10 ma x 10 ns

Magnetic field: Operates parallel to 0.7 T

Full Scale: 5 TeV jet

Min Scale: 0.1 min ionizing ~ 50 MeV

Gain: Max Charge = 200 mA x 10 ns = 2 nC, 100 p.e./GeV \rightarrow Gain = 12,000

1. Absolute:

- Radiosources, cross calibrated with beam tests
 - Cs or Co source capsules on wires in tubes (like CDF)
- Landau Distributions
 - See SPACAL reports; detailed difference $50 < E_{\mu} < 200$ GeV
- Pion Punch-thru
 - Tracker + min. ionize in BaF2 + isolation -> E pion known.
- Neutron Generator Activation, cross calibrated
 - 14 MeV D-T neutron generators ($10^{**9}/s$) (like UA-1)
 - Generator: 6 terminal metal-ceramic 5 cm dia x 25 cm long
 - 1 μ sec pulses, 1% D.F.

2. Relative:

- Light Injection on Fibers
 - Achieved $< 0.5\%$ in CLEO, Bugey Neutrino Expt., etc.
 - Long term: several months
 - Cross-calibrate in lab, on bench, & w/ radiosource
 - CDF, UA-1, UA-2: $\sim 1\%$ long term
 - lasers, plasma pinches, LED's, stable arcs, laser diodes, etc
 - [- Tunable ring laser, phase locked machine RF
 - > sub ns pulse, stability: 10^{-4}]
 - [- Inject front/rear at several wavelengths
 - > rad damage, separate fibers from PMT stability]
- Symmetries of Energy Deposition
 - Towers of constant $\pm\eta$ (a ring of towers): same Energy.
 - Sets of towers of constant ϕ : same $\delta E/\delta\eta$ (Energy profile)
 - Compute Entropy; anneal
 - Use tracker to correlate with absolute $d^3E/dtd\phi d\eta$.

A SIMPLE METHOD FOR THE STABILIZATION OF SCINTILLATION DETECTORS

M. BÖTTCHER, W.D. BREWER, and E. KLEIN

Fachbereich Physik, Freie Universität Berlin, W. Germany

Received 10 May 1982

We report on the design and operation of a gain-stabilized NaI(Tl) scintillation spectrometer for use in stray magnetic fields utilizing a blue light-emitting diode as reference source. The spectrometer is simple, compact, and robust and the stabilization can be constructed at modest additional cost. It is easy to operate, and yields long-term gain stability of better than $\pm 0.2\%$, as well as greatly improved operation in stray magnetic fields.

Nuclear Instruments and Methods in Physics Research A279 (1989) 73-76
North-Holland, Amsterdam

(~ 300 p.p.t.) 73

PERFORMANCE OF A LIGHT FLASHER SYSTEM FOR A PROTOTYPE CsI CALORIMETER AND THE DESIGN FOR CLEO II

Yuichi KUBOTA and Chris O'GRADY *

Cornell University, Ithaca, New York 14853, USA

→ Have now reported
long term $\pm 0.5\%$

A study of the light calibration system installed in a prototype CsI shower counter is reported. The system is able to observe a change of less than 0.5% in the sensitivity of the detector.

Nuclear Instruments and Methods in Physics Research A234 (1985) 517-520
North-Holland, Amsterdam

517

GAIN STABILIZATION OF PHOTOTUBES USING A LED DIODE SCHEME *

L. HOLM, H.W. FIELDING and G.C. NEILSON

Nuclear Research Centre, Physics Department, University of Alberta, Edmonton, Canada, T6G 2N5

Received 13 July 1984

The performance of LED diode gain stabilization schemes for RCA 4522 phototubes is evaluated. Under normal experimental conditions the use of a green Litronix GL56 LED and an ORTEC surface barrier detector provide the best results. An overall gain stabilization of $\pm 0.5\%$ over several months has been achieved.

LASER/FIBER OPTIC CALIBRATION SYSTEM

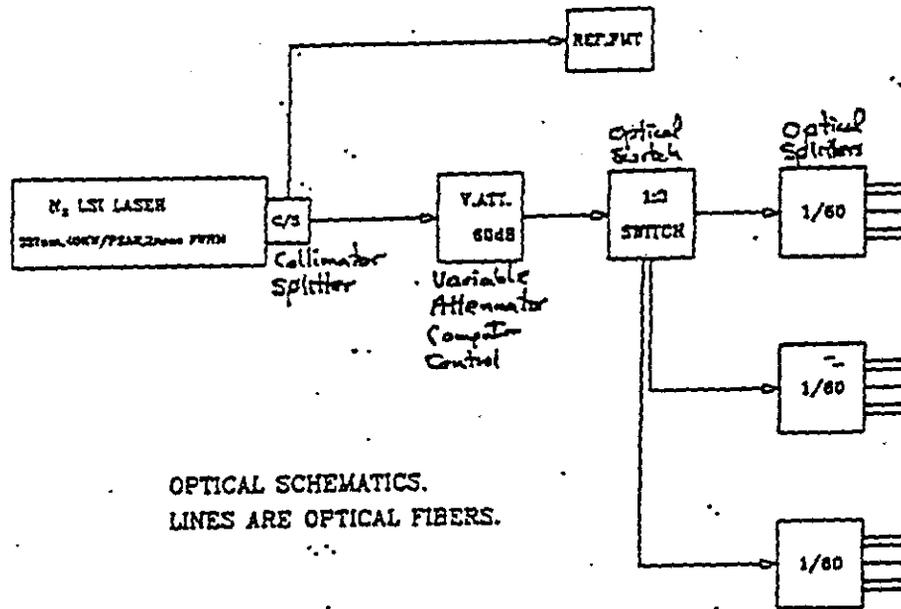


Fig. 2a: Schematic of full optical pulser system for a large scale calorimeter system.

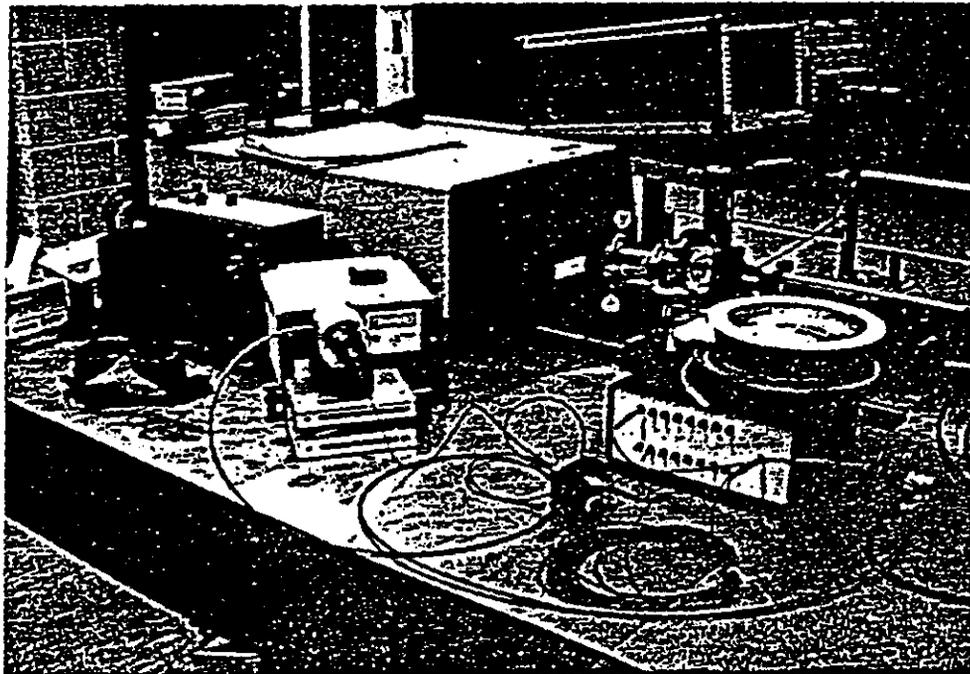
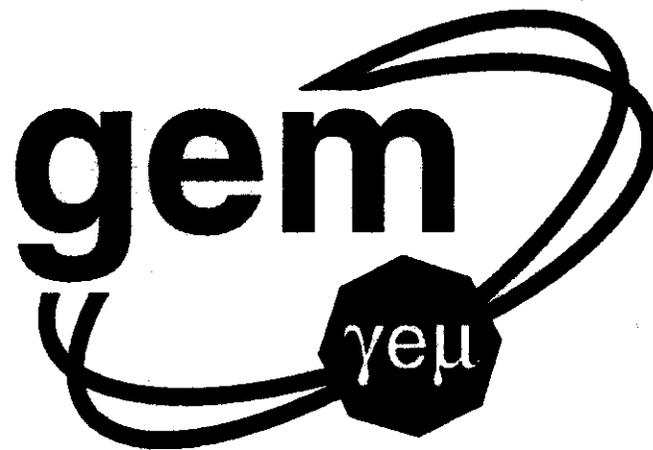


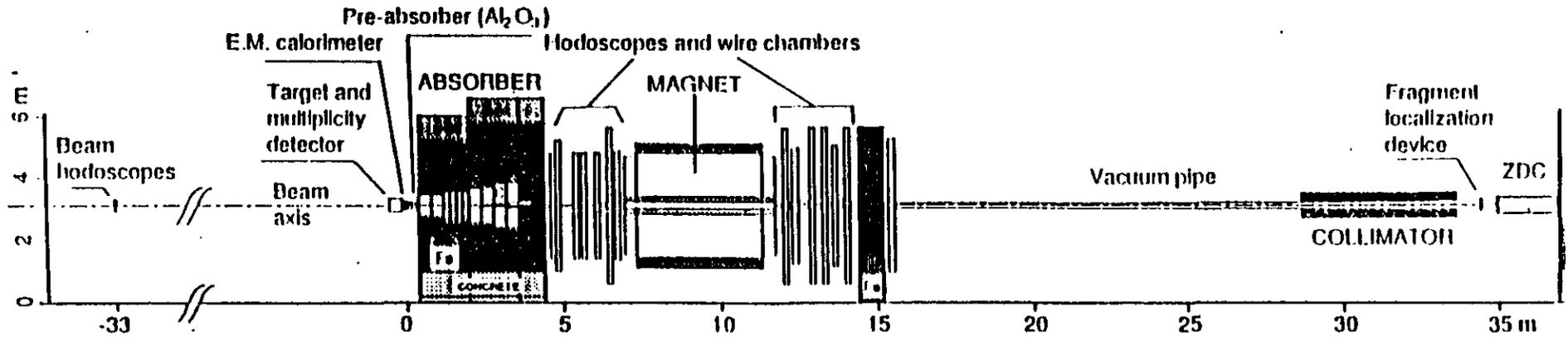
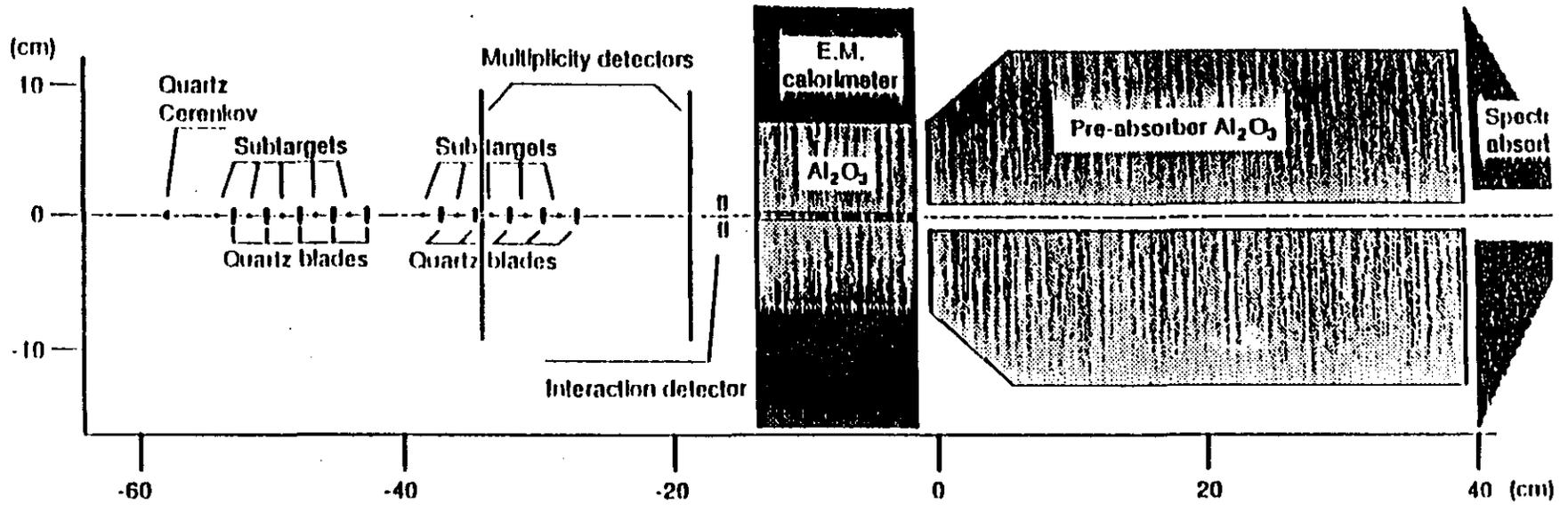
Fig. 2b: Photograph of early test assembly of the 16 channel light pulser calibration system.



Presentation by:

Philippe Gorodetzky

III.5: Vue générale de la nouvelle expérience.



REQUIREMENTS FOR NA38 with Pb Beam

1) Fast response

Intensity is $1.0 \cdot 10^7$ ions/sec

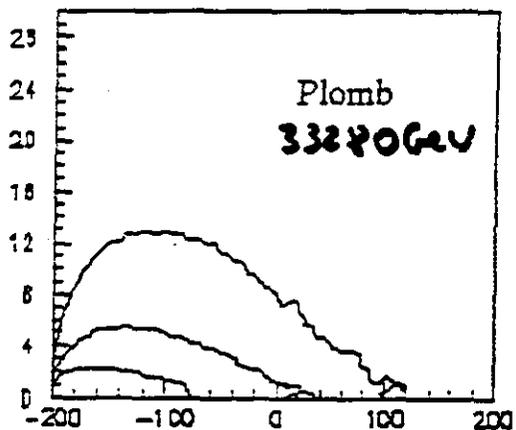
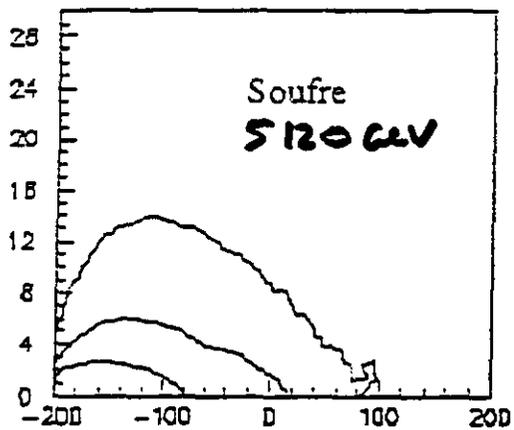
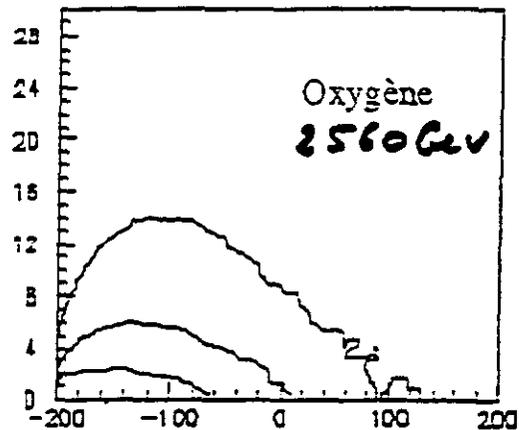
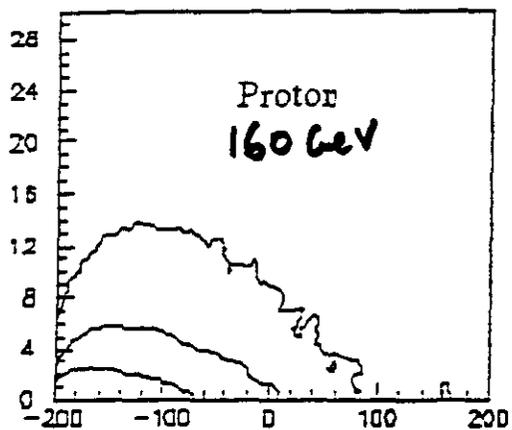
⇒ pulse should be back
to zero before 20 ns

2) Radiations

shall suffer 2-5 Grads (10^7 Gy)

3) Resolution in hadronic mode better than 10%

4) Ease of construction



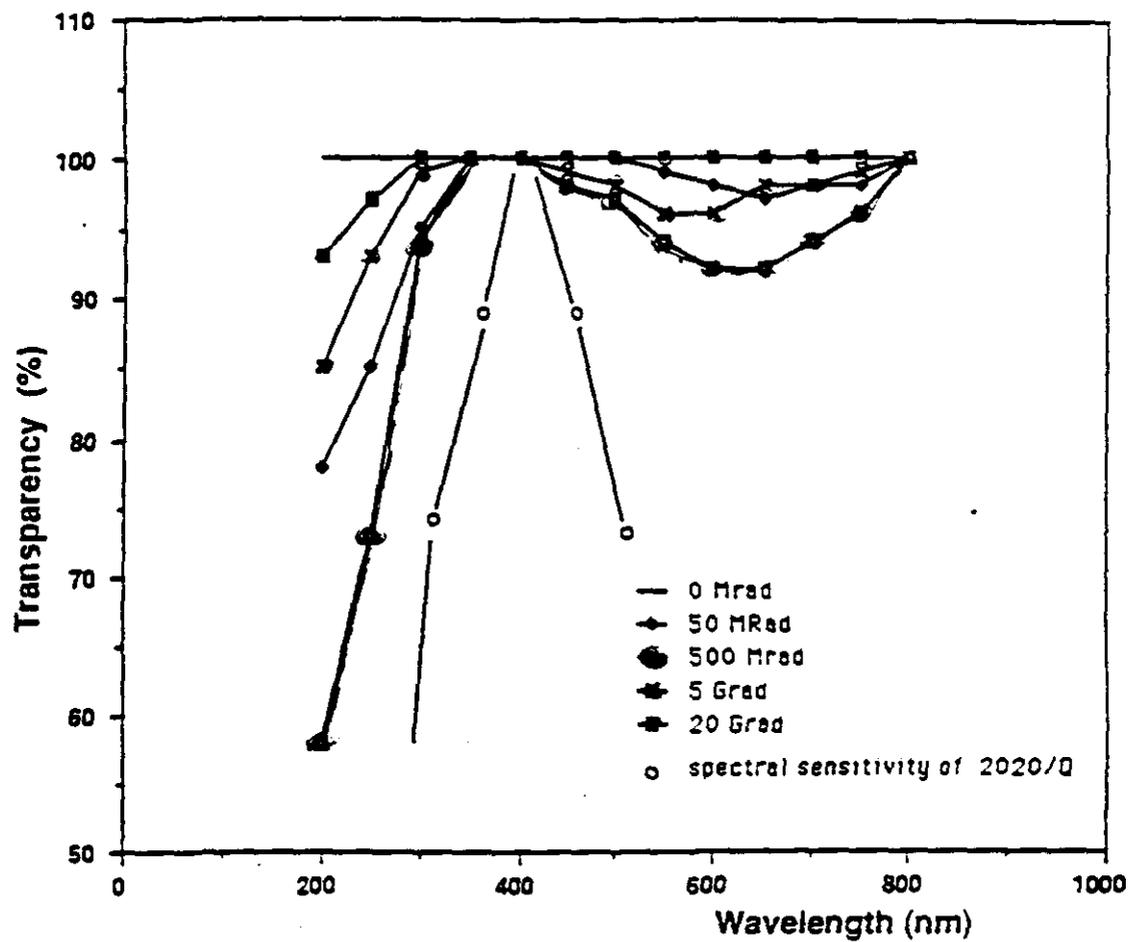
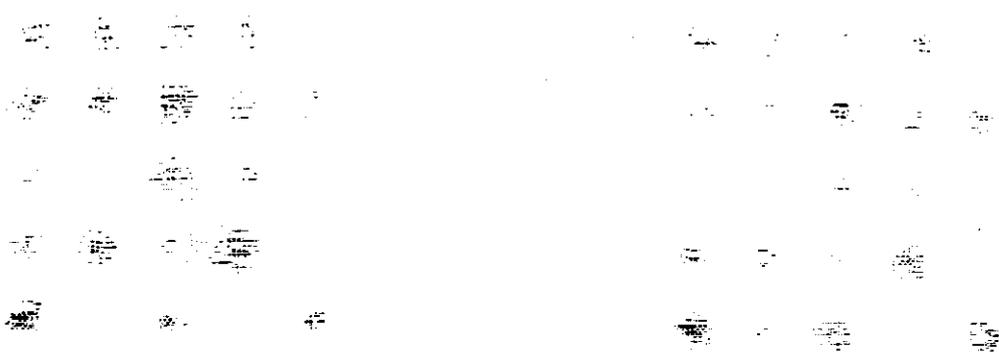


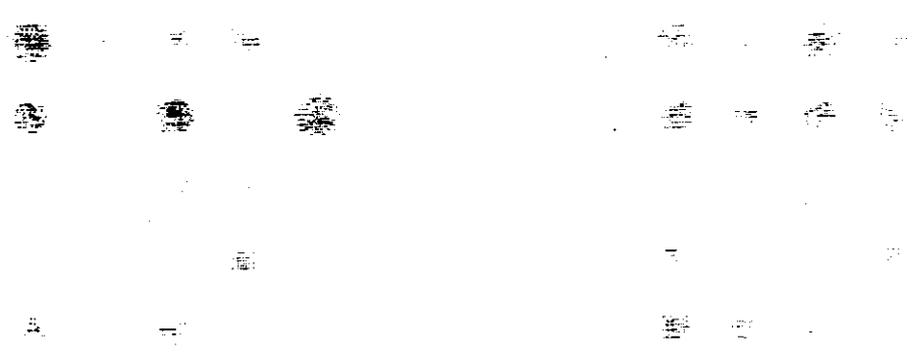
Figure 6 : Transparency of a 2 mm thick Suprasil quartz disc after different doses of irradiation. The spectral sensitivity of the 2020 Q photomultiplier shows the interesting fact that this wavelength range is the most insensitive to radiation.

Duke's transparent



8:10:32

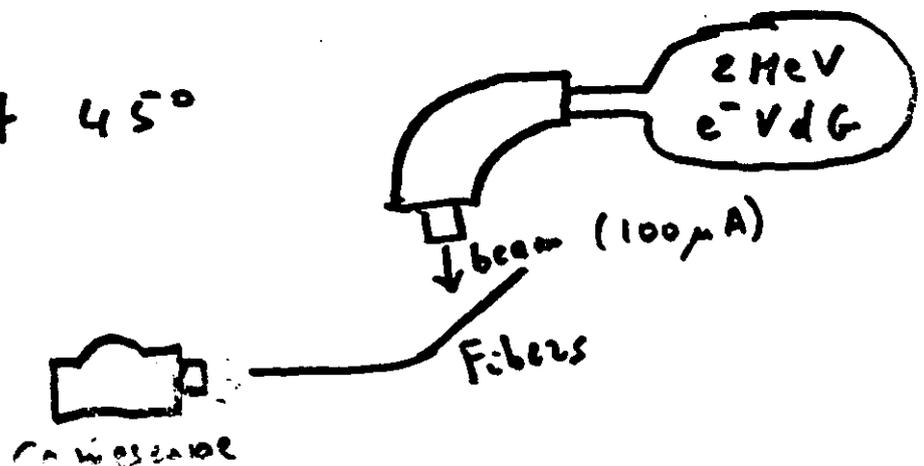
9:00:09



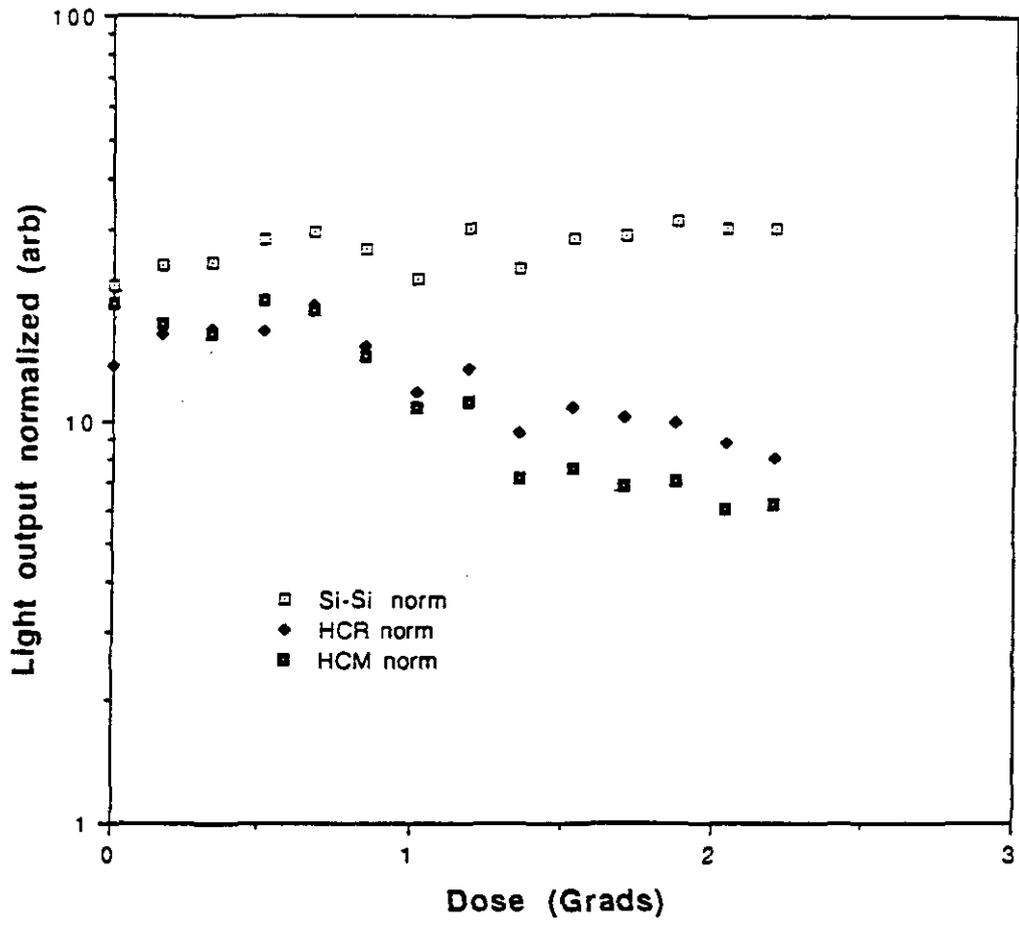
9:50:21

10:50:09

Fibers at 45°



Quartz fiber irradiation with electrons



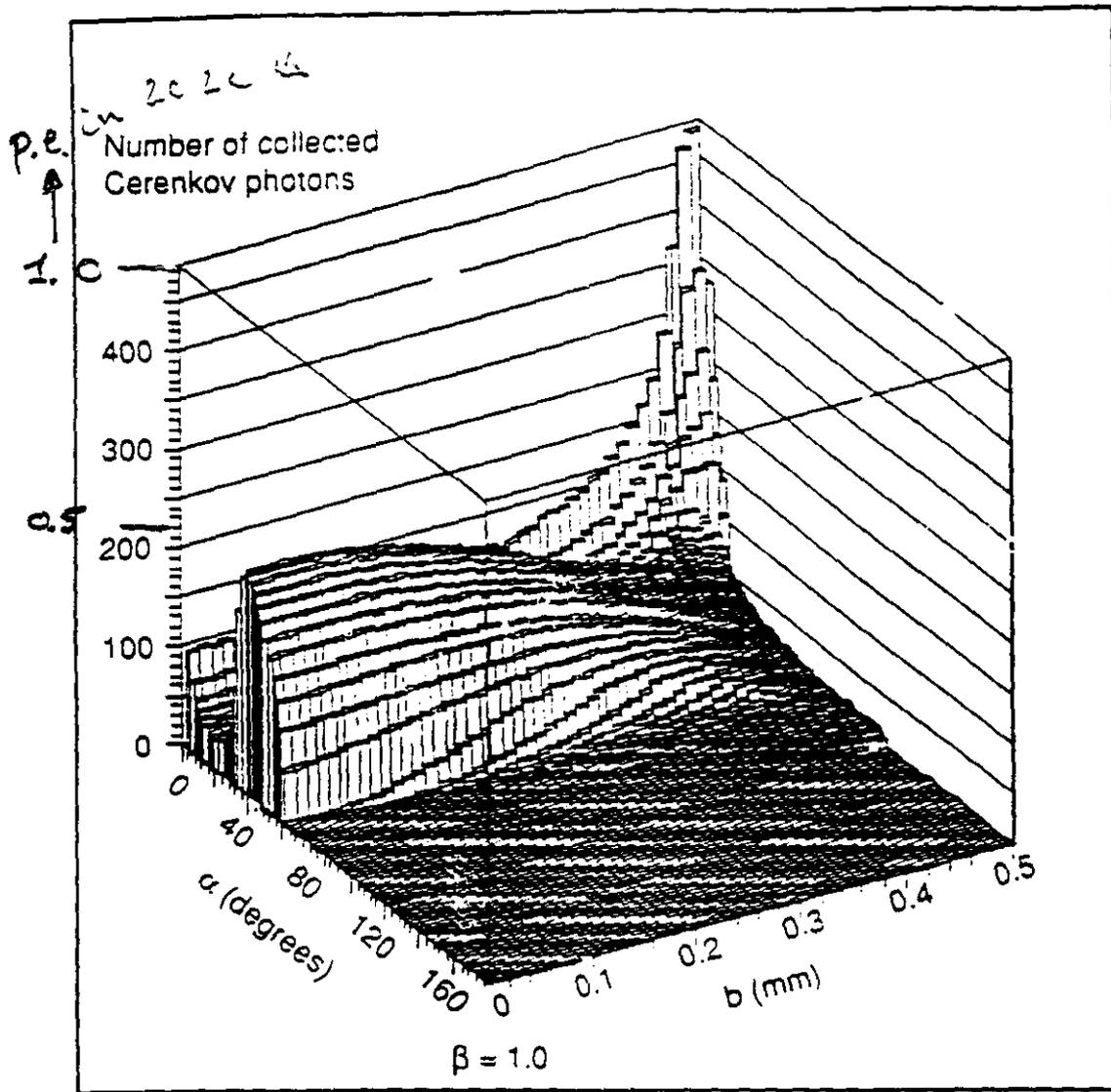
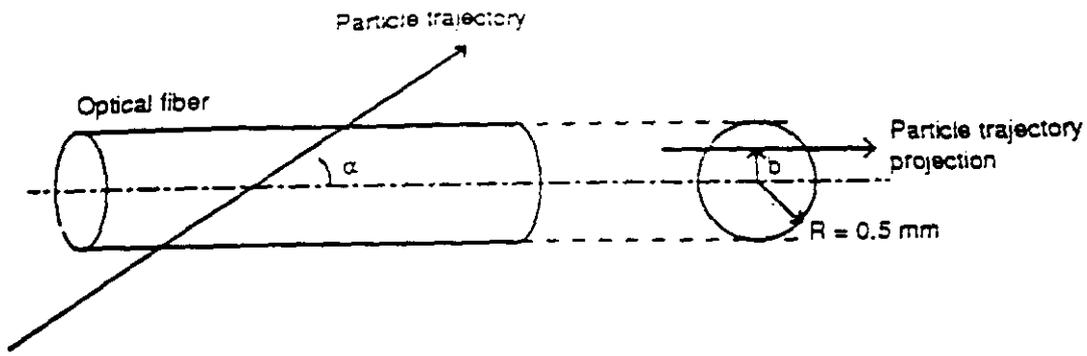


Figure 2 : Top) Geometrical parameters of an optical fiber - particle trajectory system. Bottom) Number of photons produced by Čerenkov effect and collected at the extremity of a fiber (1 mm diameter and numerical aperture of 0.37) intercepted by a charged particle ($Z=1$, $\beta = 1$).

$\beta = 1$, $\theta = 1$ at $b = 0$ through the center of a fiber, 1 mm thick gives 0.44 p.e.

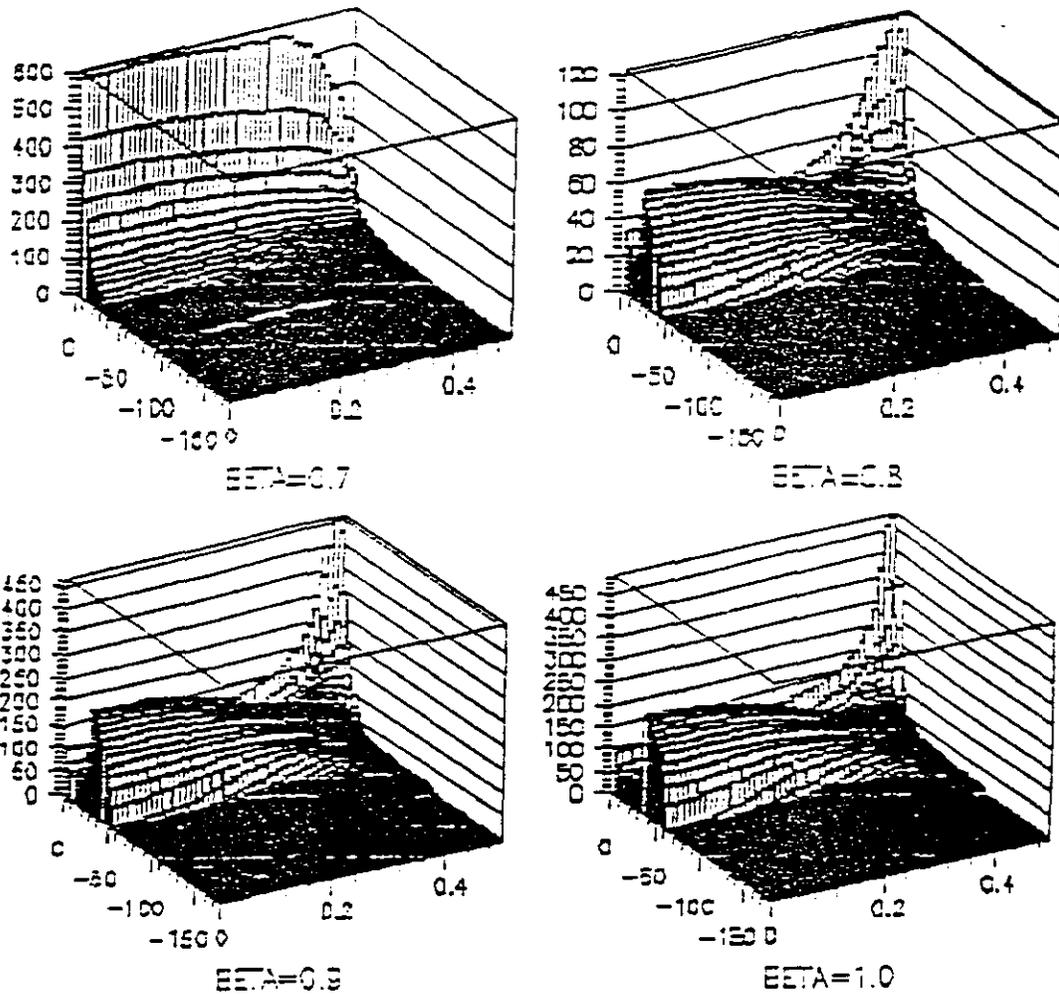


Fig V.22: Nombres de photons (quelle que soit leur longueur d'onde) arrivant en bout de fibre en fonction de l'angle et du "paramètre b" exprimés en mm pour les 4 intervalles de β .

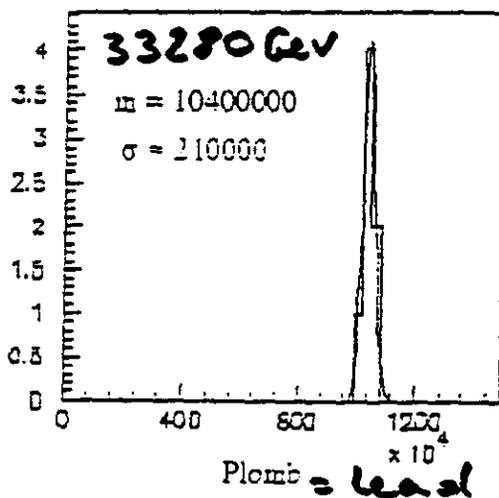
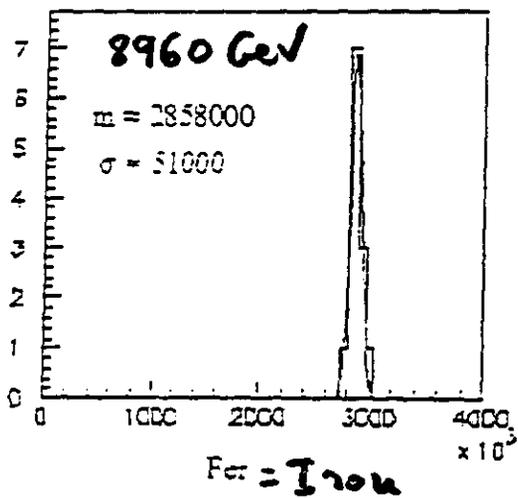
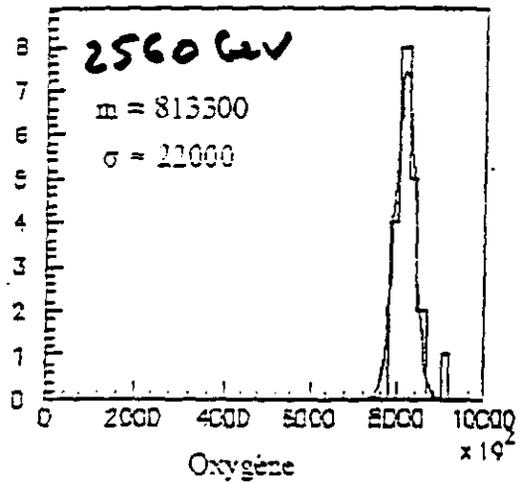
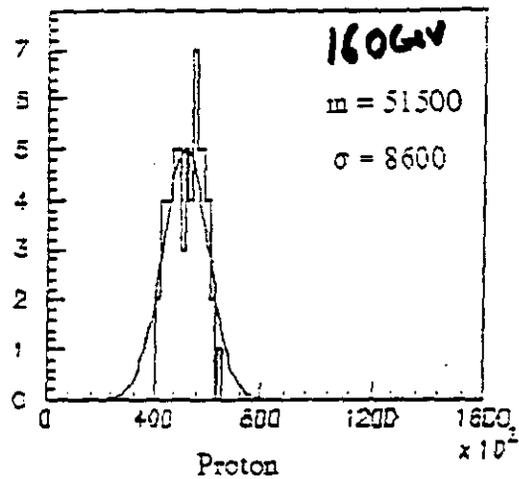
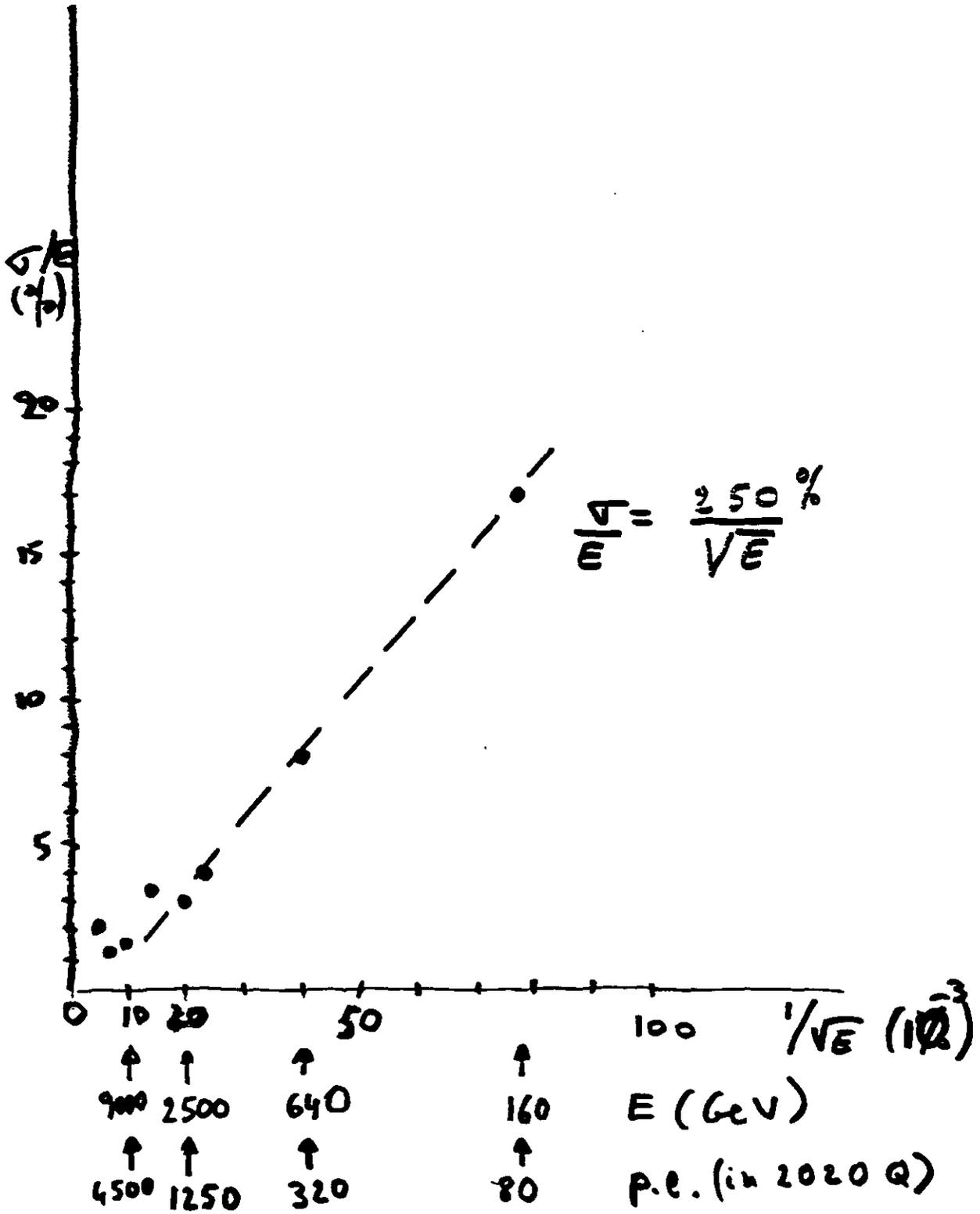


Fig V.29.a: Résolution en proton, oxygène, fer et plomb.



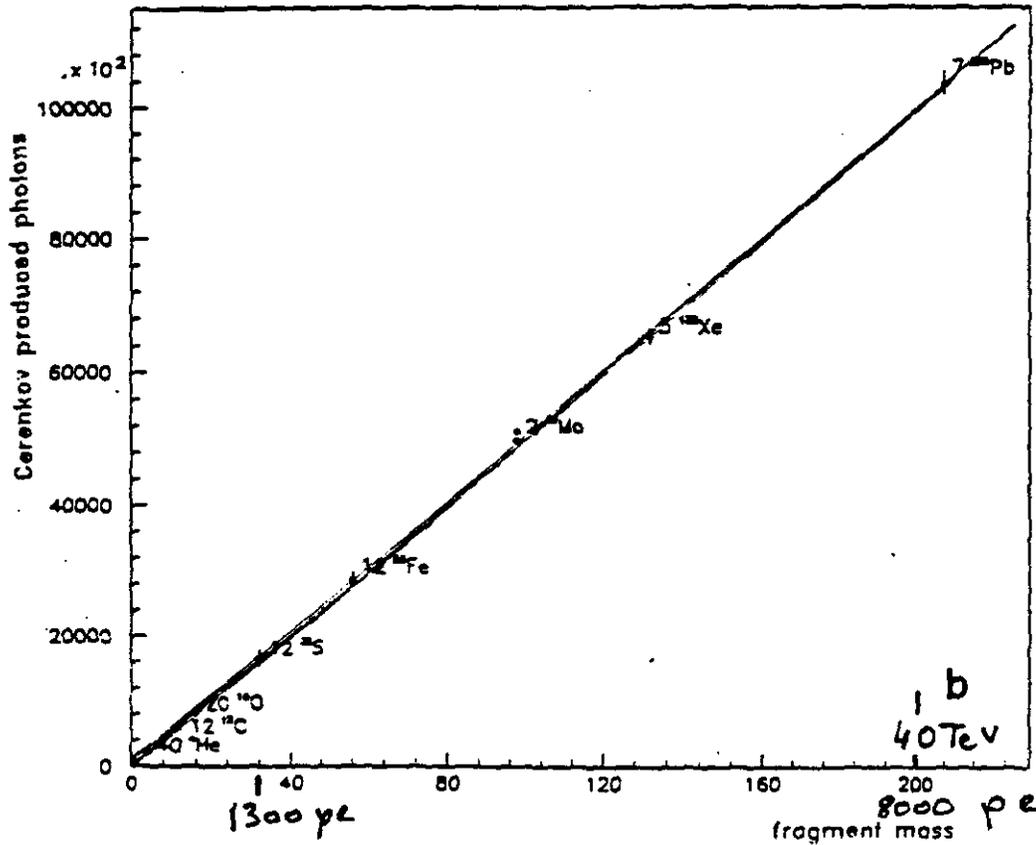
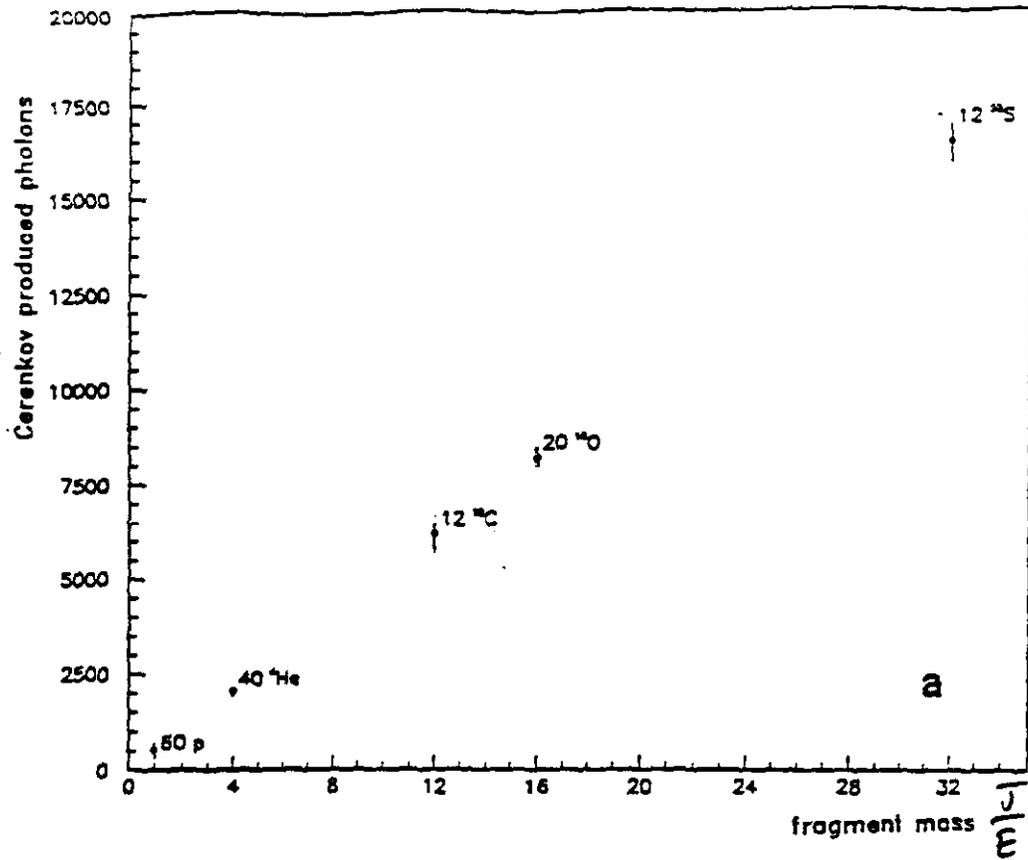
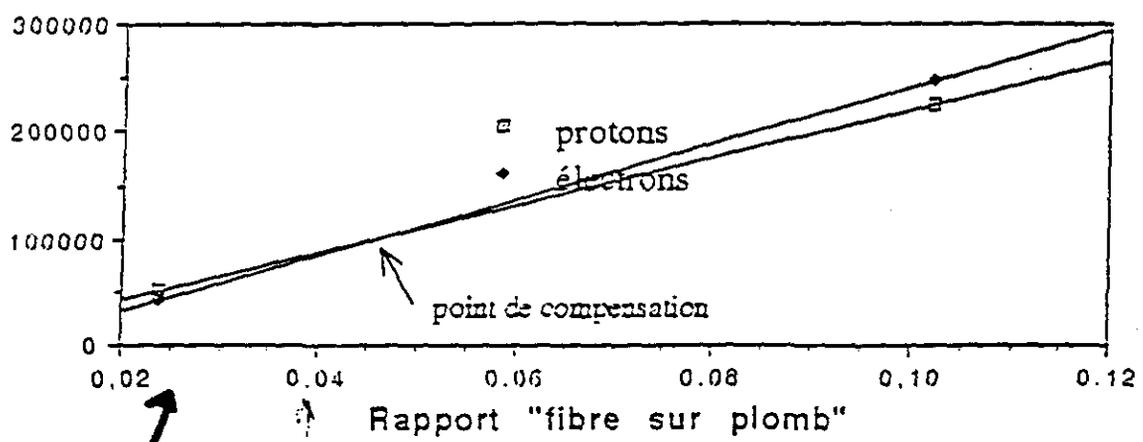


Figure 4 : Linearity of the ZDC. Figure on the top is a magnification of the low mass region. Near each point is indicated the number of events calculated by the simulation program. The error bars correspond to the FWHM energy resolution as deduced from a least squares fit. See text for more details.



"our" ratio

minimum

$$E_{e^-} = E_p = 160 \text{ GeV}$$

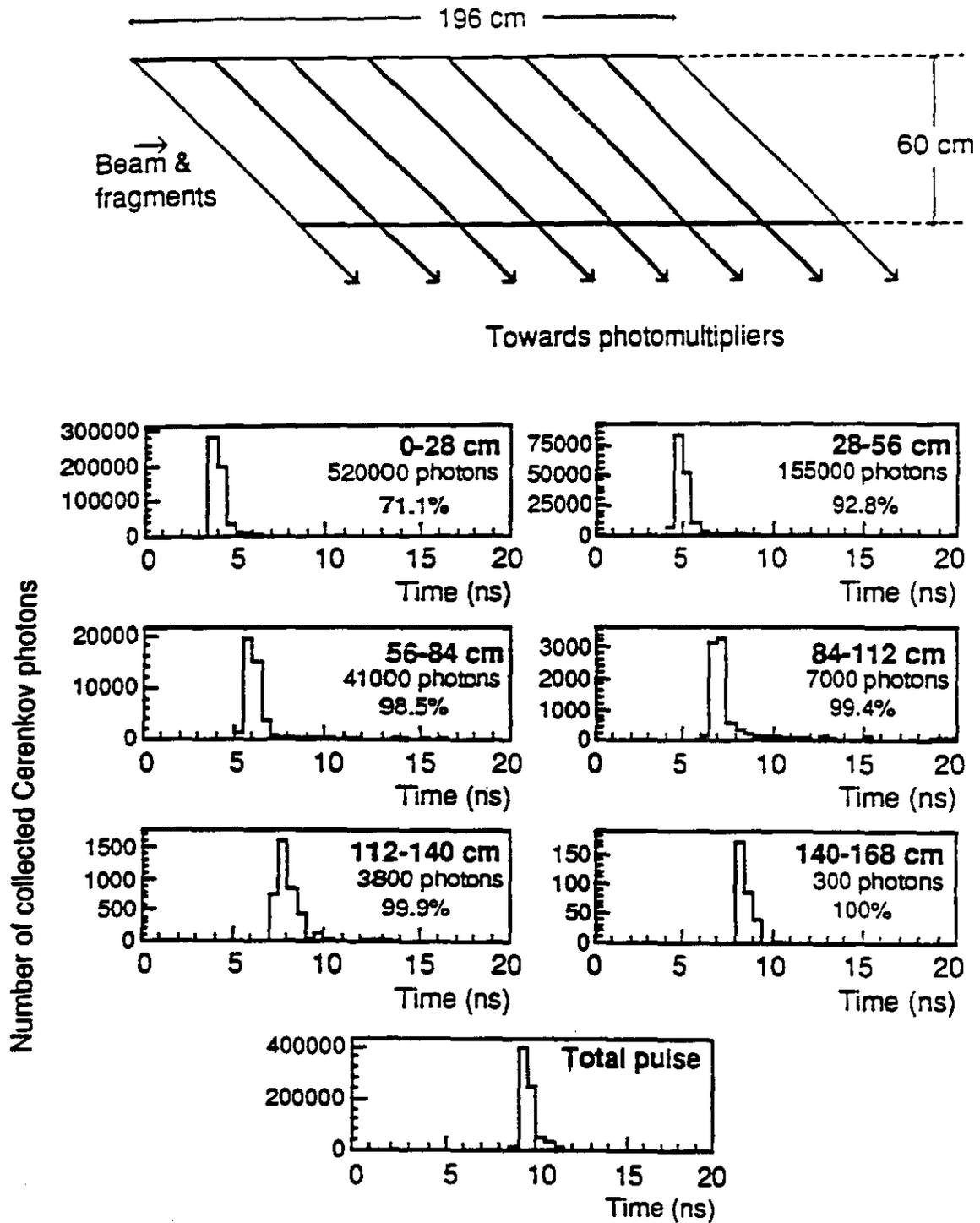


Figure 5 : Top) Schematic view showing the seven 28 cm thick slices along the Z-axis. Bottom) The anode pulses from the Čerenkov light from the different slices collected by the different photomultipliers as a function of time for an Oxygen ion of 160 GeV/nucleon. Pulses of adjacent slices are shifted by one nanosecond due to the shower development along the z-axis with a value of β close to 1. The full shower for this event is contained inside 6 slices. The last pulse is the sum of the six pulses, each of them being delayed by one nanosecond relative to the next one. It is the useful pulse for the experiment.

P. GORODETZKY
D. LAZIC

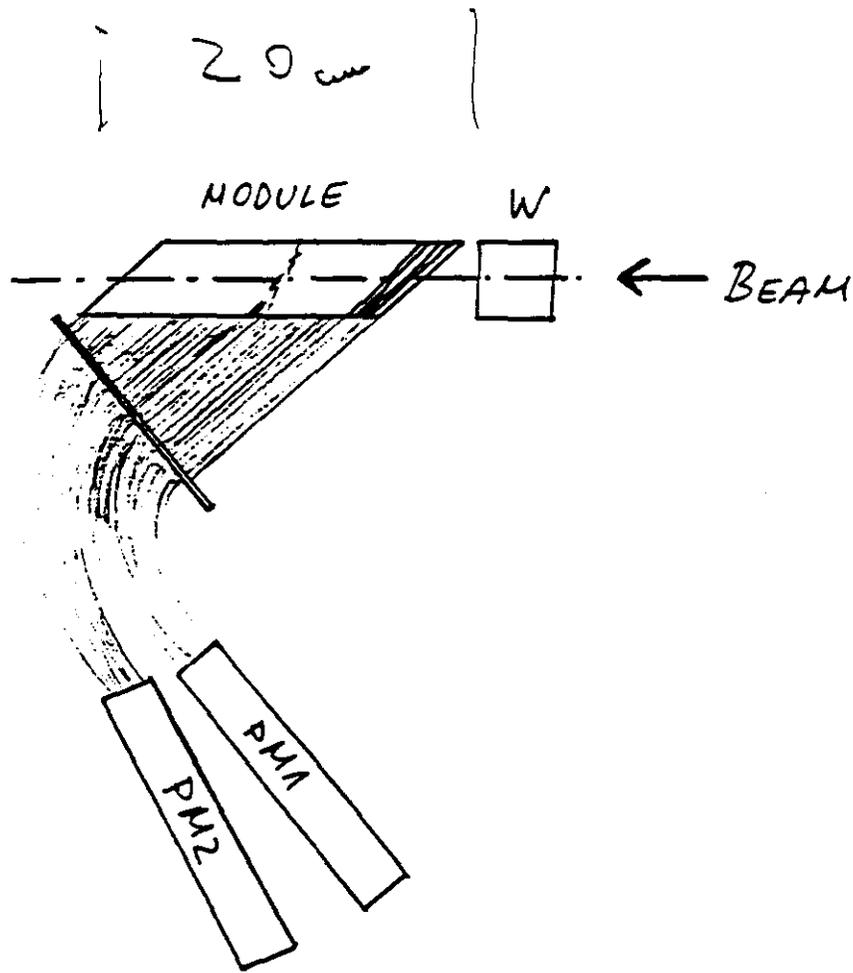
SILICAL

CALORIMETRY WITH SILI(CA) FIBERS

Tests performed at CERN on the small prototype with S^{32} at 200 GeV/n
and electrons at 10 and 30 Gev

- PRELIMINARY RESULTS -

October 1991



DIMENSIONS OF THE MODULE: $5 \times 5 \times 20$ cm

THICKNESS OF TUNGSTEN BLOCK: 0, 4, 8 cm

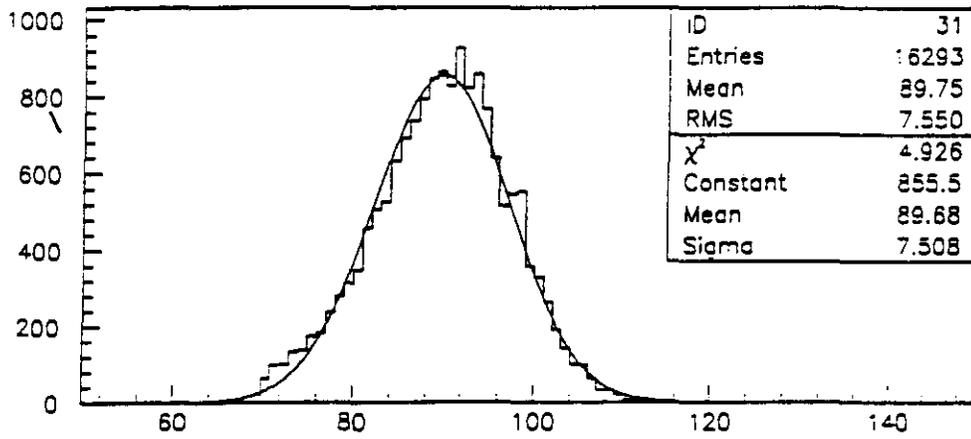
1474 FIBERS 50 cm LONG

DISTANCE BETWEEN FIBERS 2.22, $V_{PB} : V_{Fib.} = 4:1$

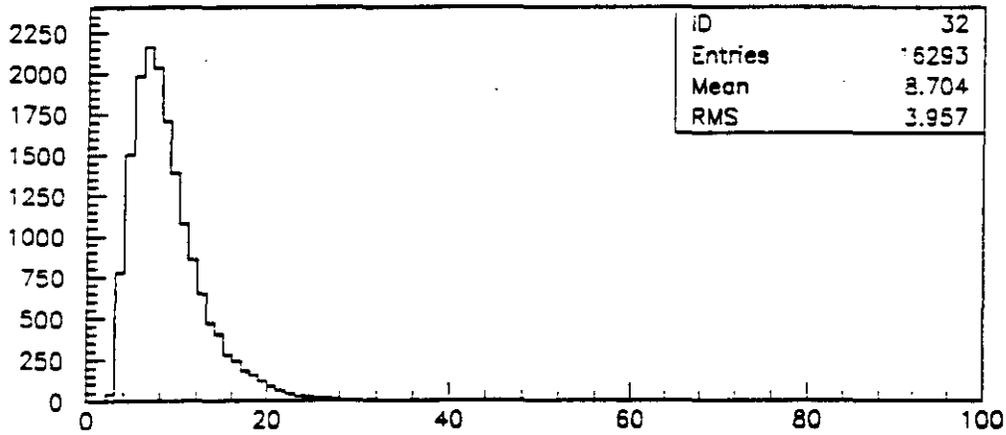
INTERACTION LENGTHS FOR S^{32} :

- IN W $\lambda_{int} = 4$ cm

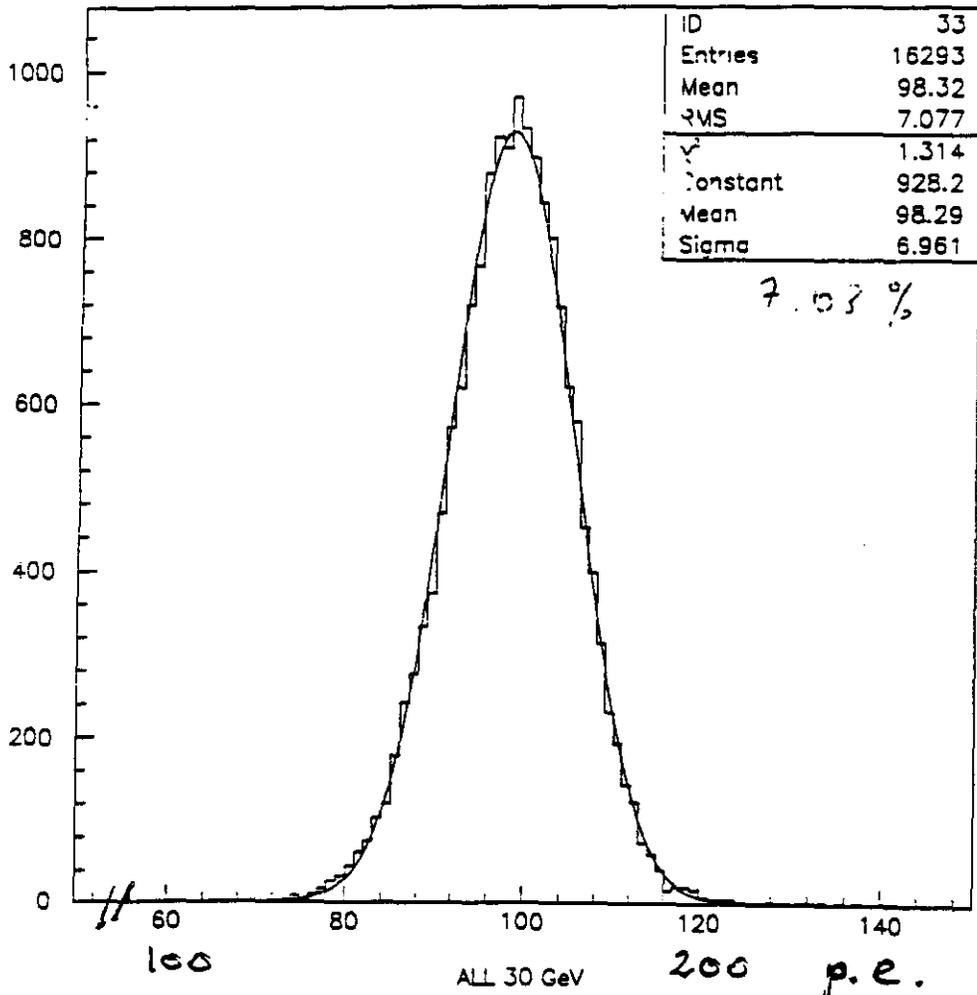
- IN MODULE $\lambda_{int} = 9.1$ cm (including quartz fibers)



ADC1 30 GeV



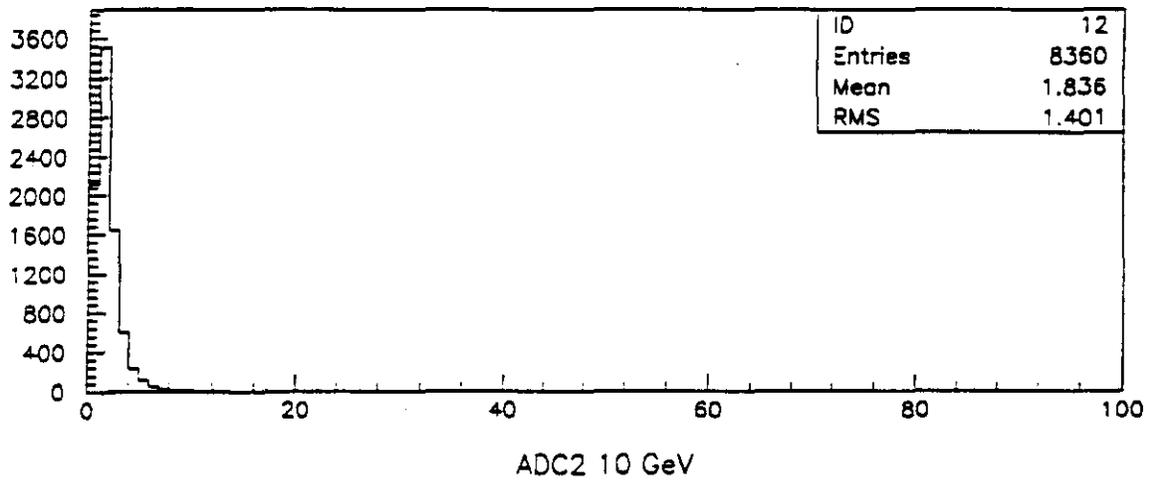
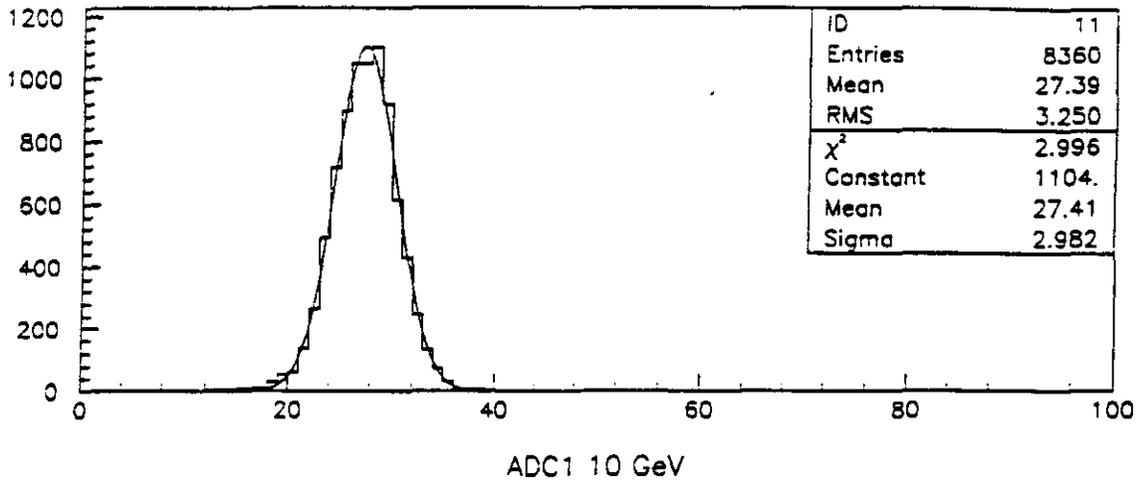
ADC2 30 GeV

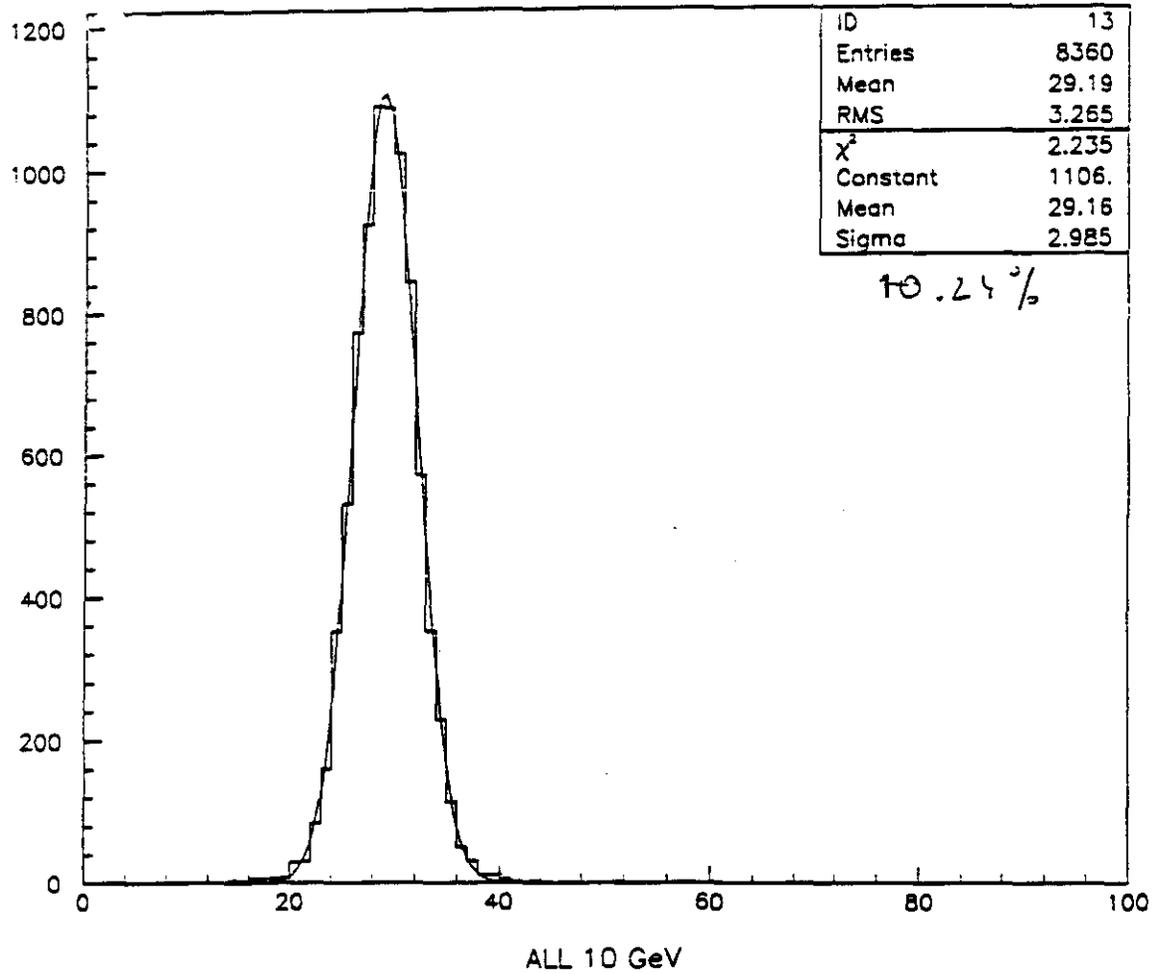


$$\Rightarrow \frac{\sigma}{E} = \frac{23.6\%}{\sqrt{E}} + 2.77\%$$

$$\frac{\sigma}{E} = \frac{a}{\sqrt{E}} + b$$

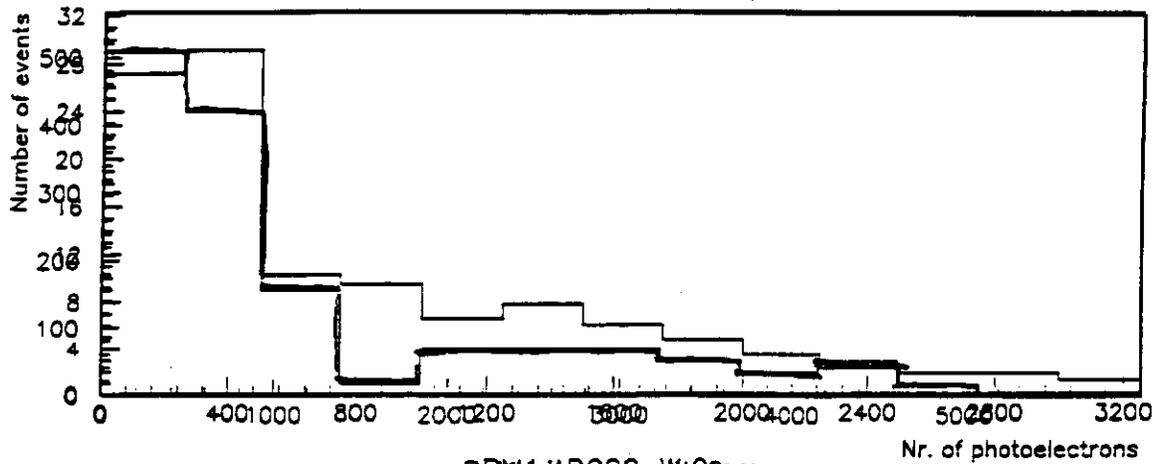
~~100~~ 6 p.e. / GeV





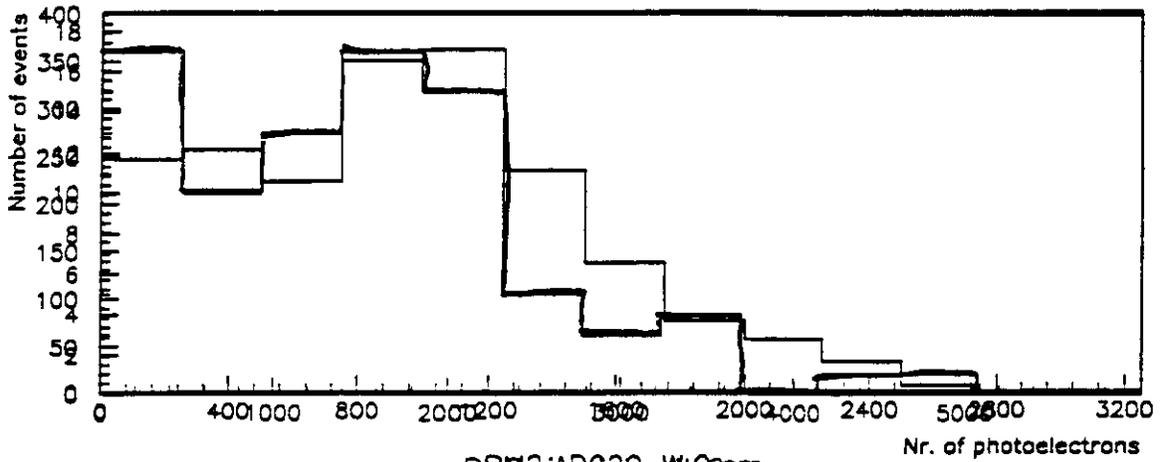
SIMULATED

Sulphur tests - MEASURED



pe

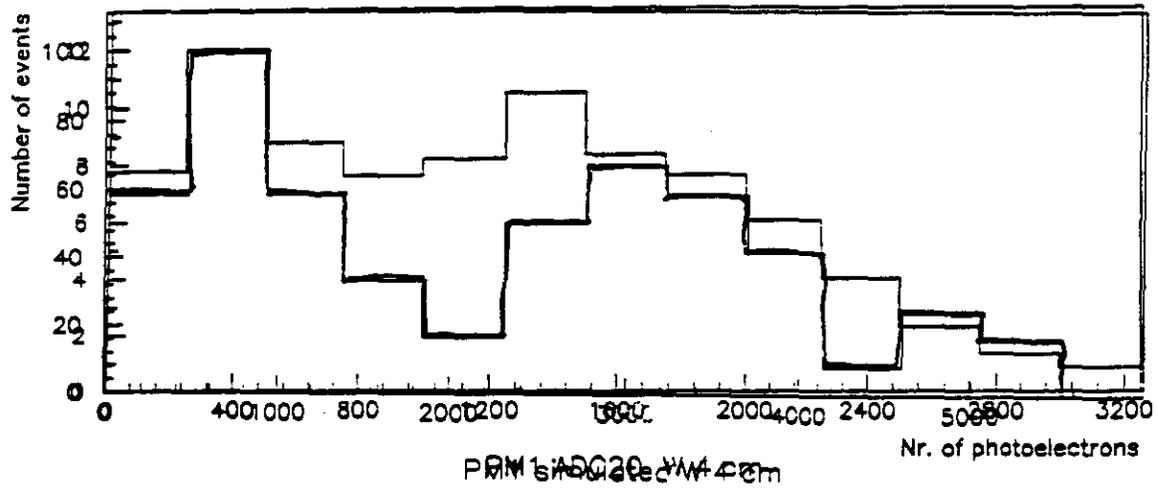
PRM 21ADG20dW00com



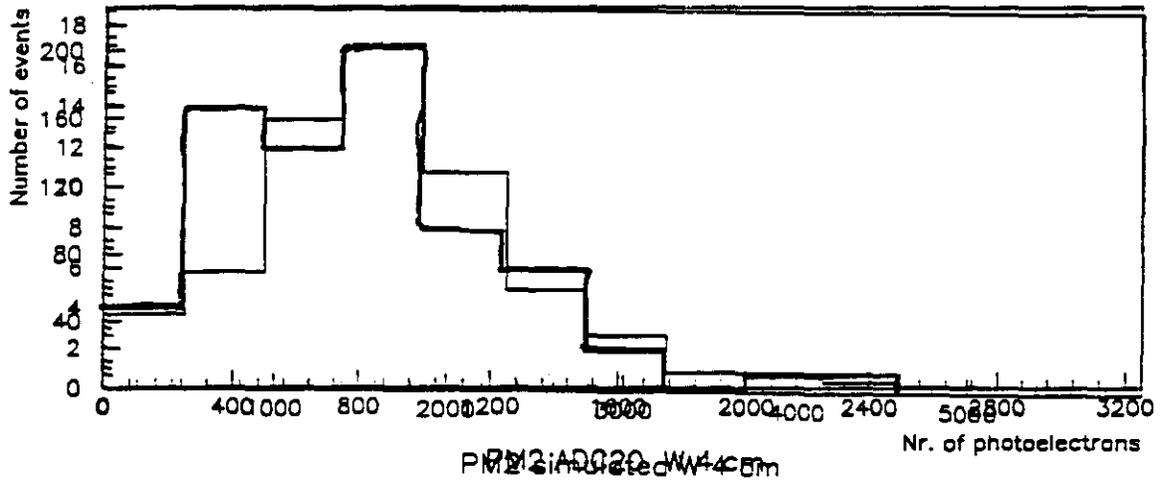
PRM 21ADG20dW00com

SIMULATION

Sulphur tests - MEASURED

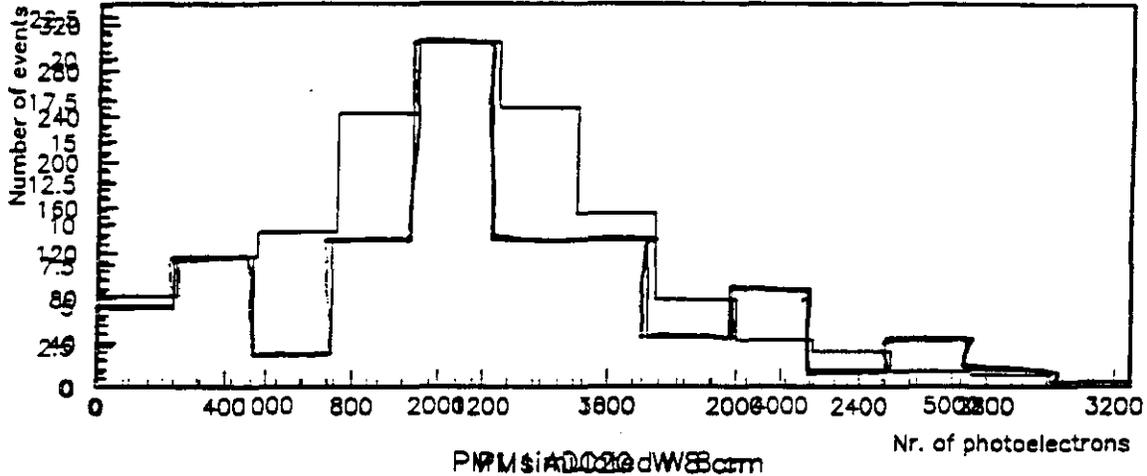


p.e.

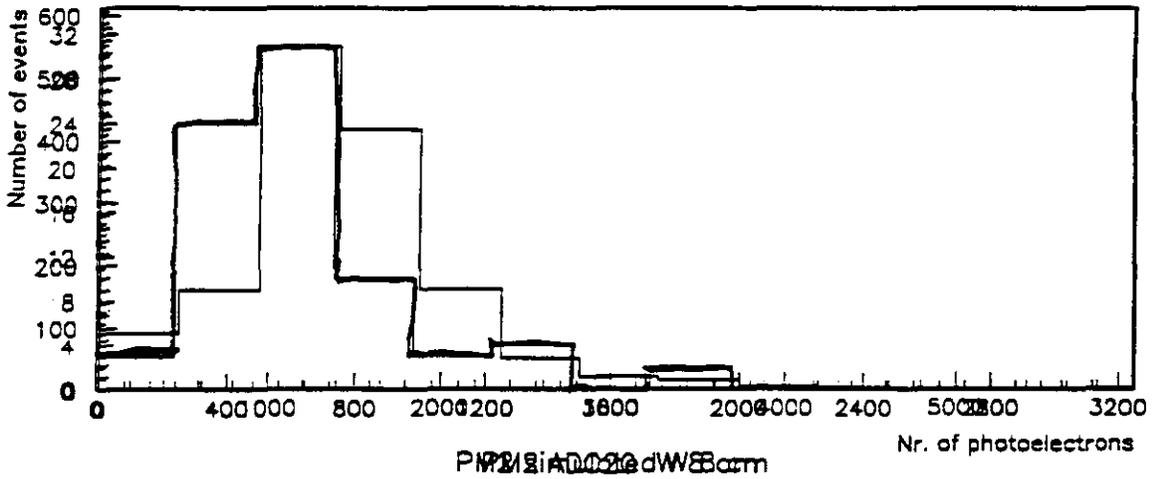


SIMULATED

Sulphur tests - MEASURED



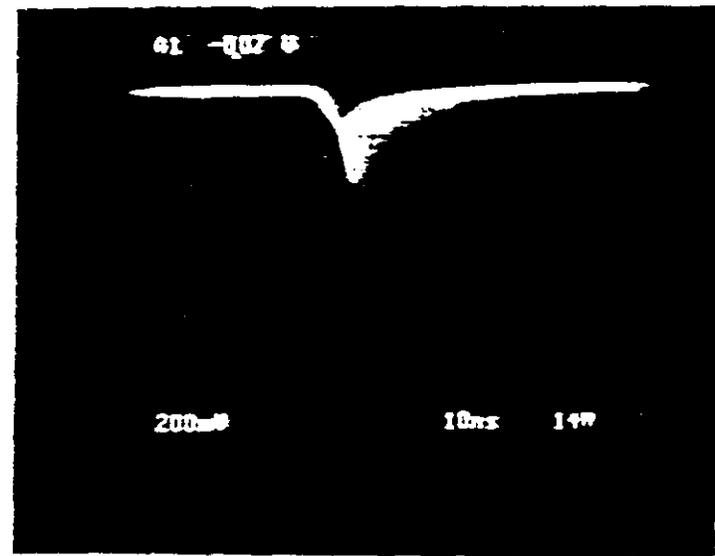
200
1150
2000
P.e.



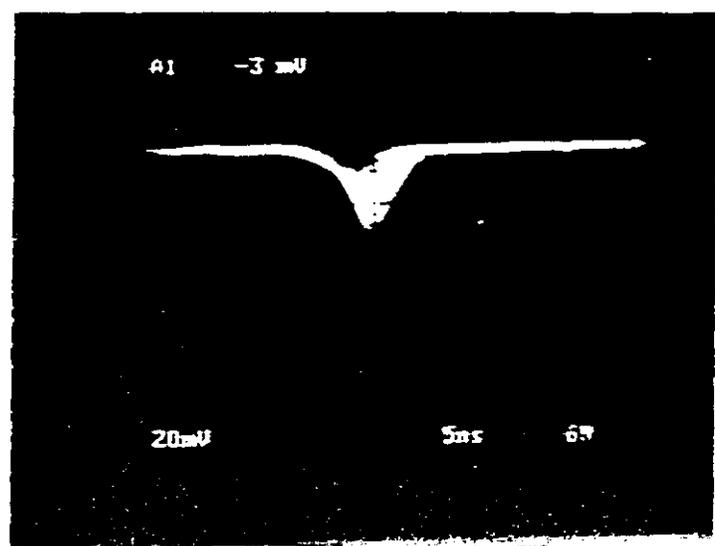
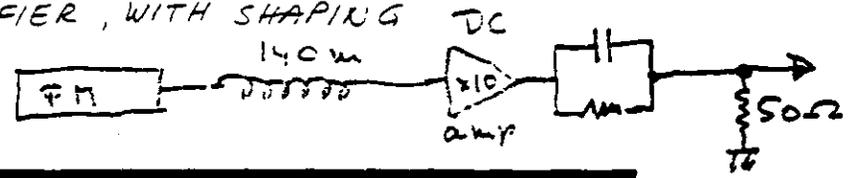
1020

32 TEST Run W PULSE SHAPES

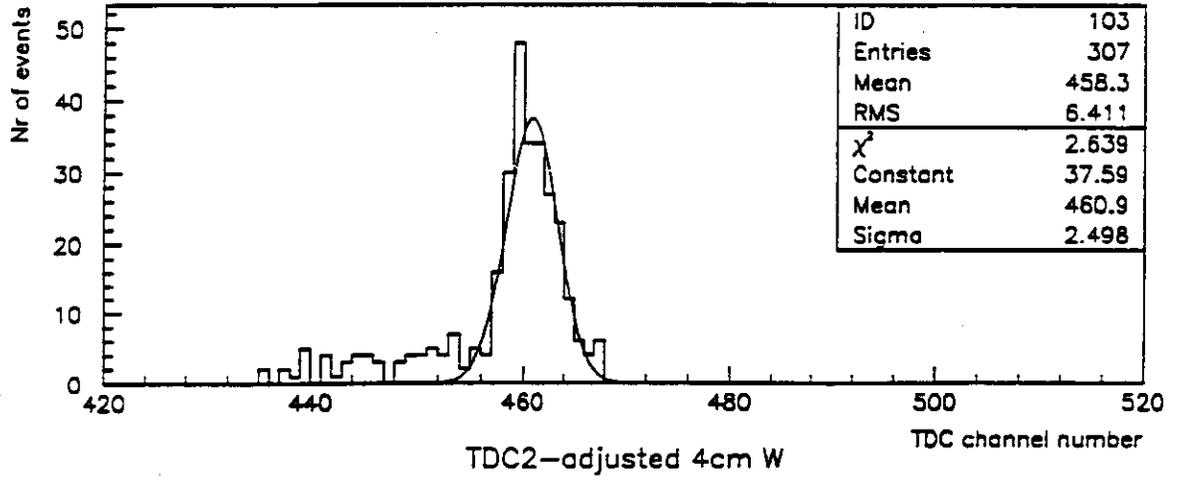
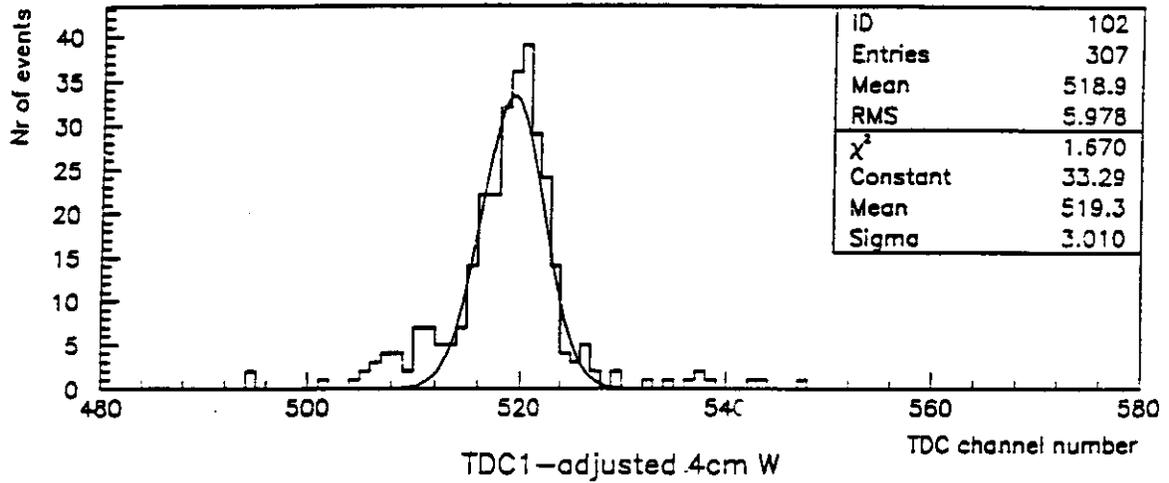
PMA, TWO AMPLIFIERS, NO SHAPING (140 m CABLE)



PMA, ONE AMPLIFIER, WITH SHAPING

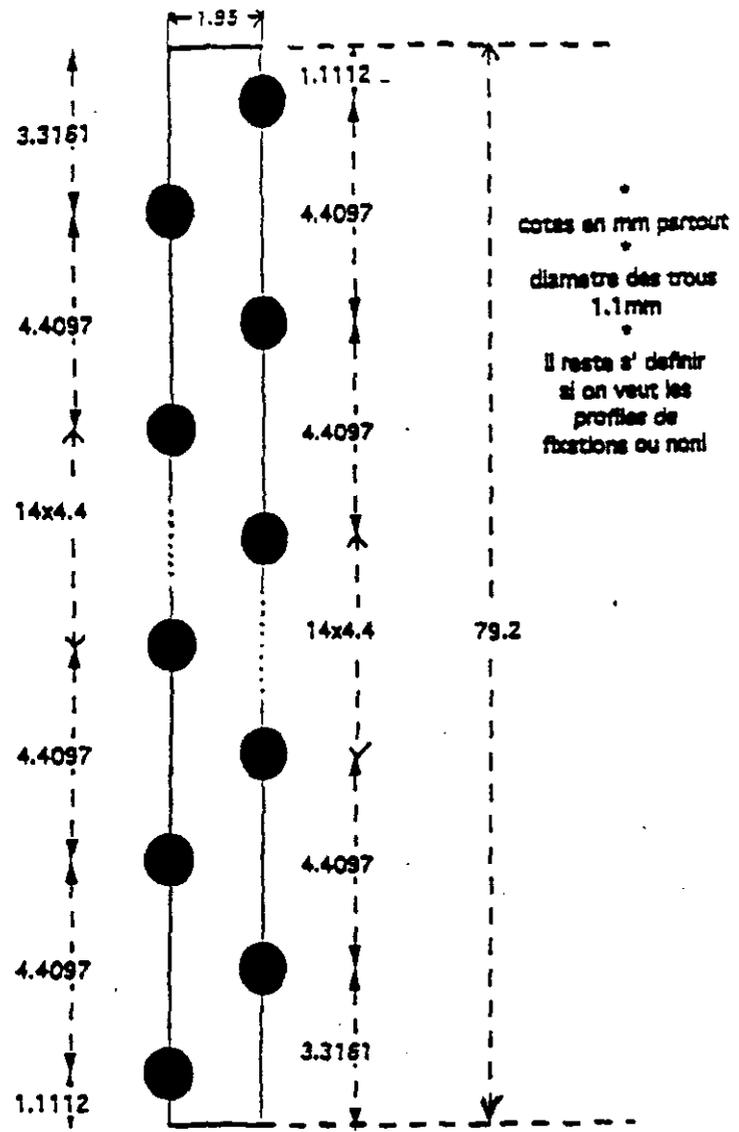


TIME RESOLUTION AFTER THE TRANSFORMATION:

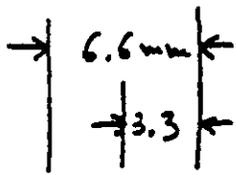


$$\sigma = 250 - 300 \text{ ps}$$

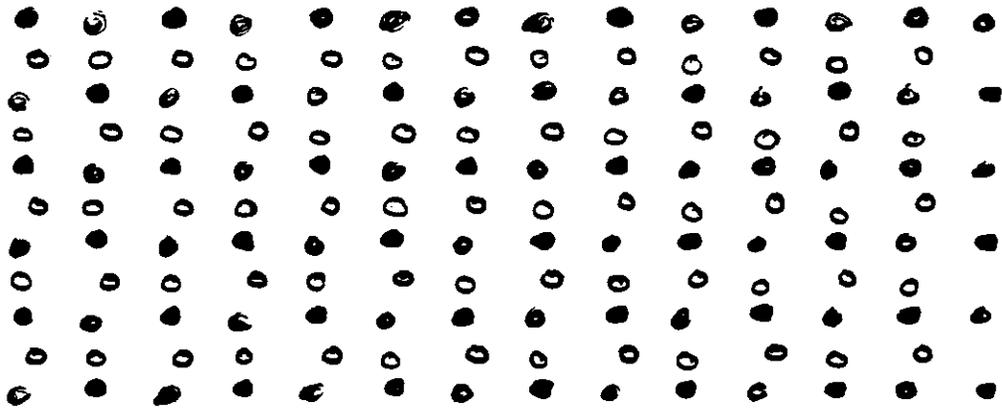
LAMIN PLATES (ORDERED) :

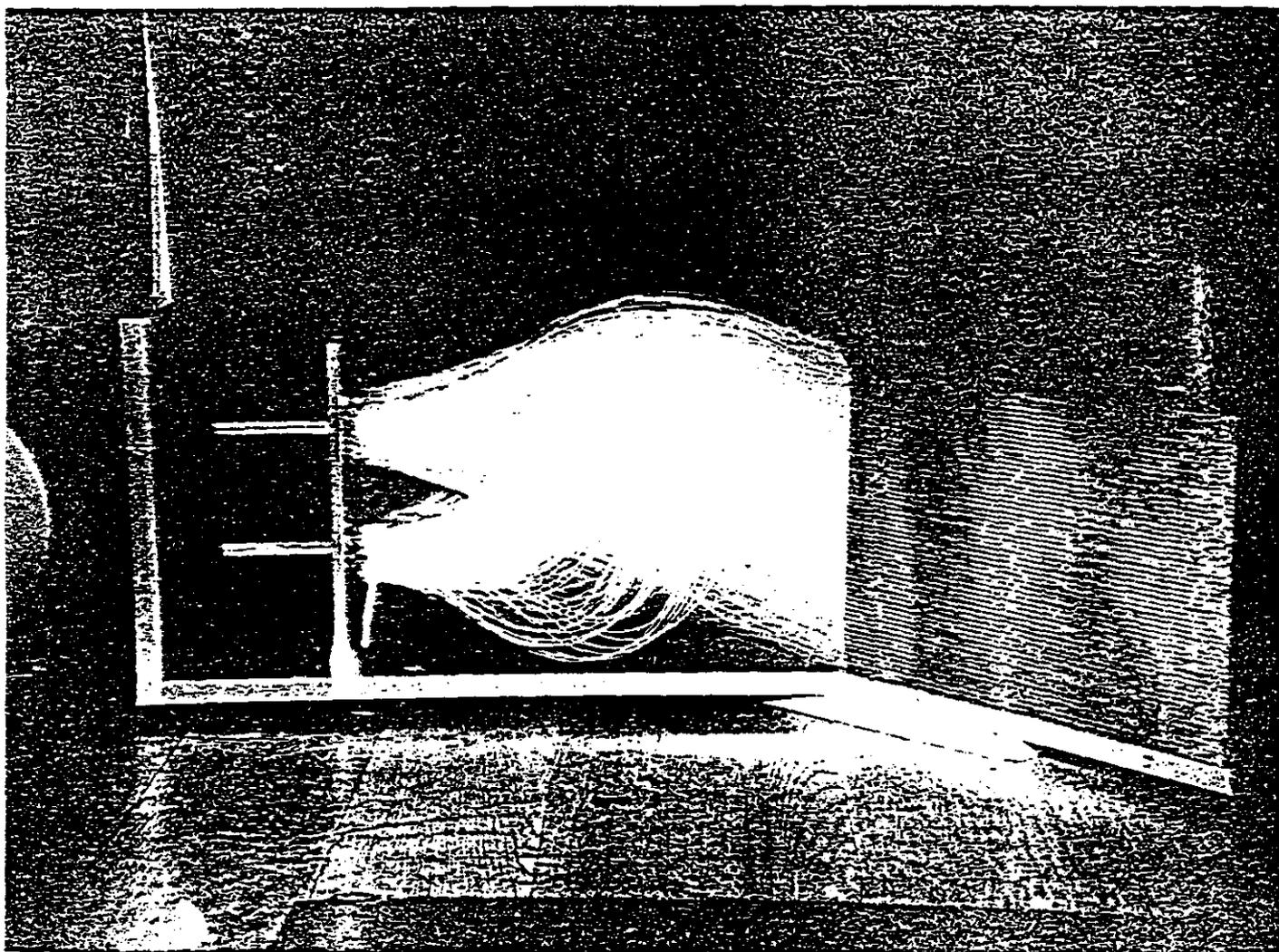


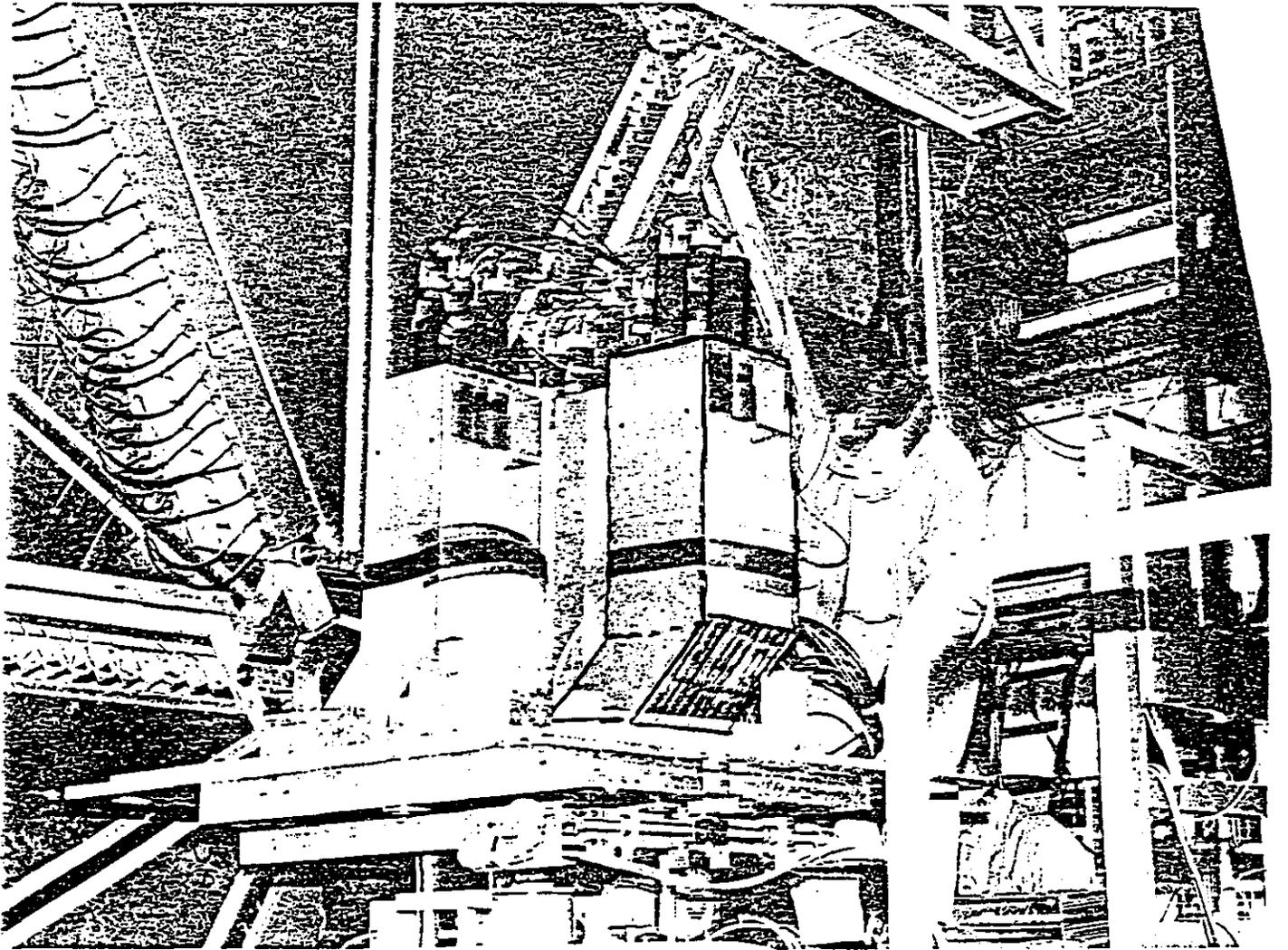
fibers and
 2 fiber is
 (see triangles)
 on side and

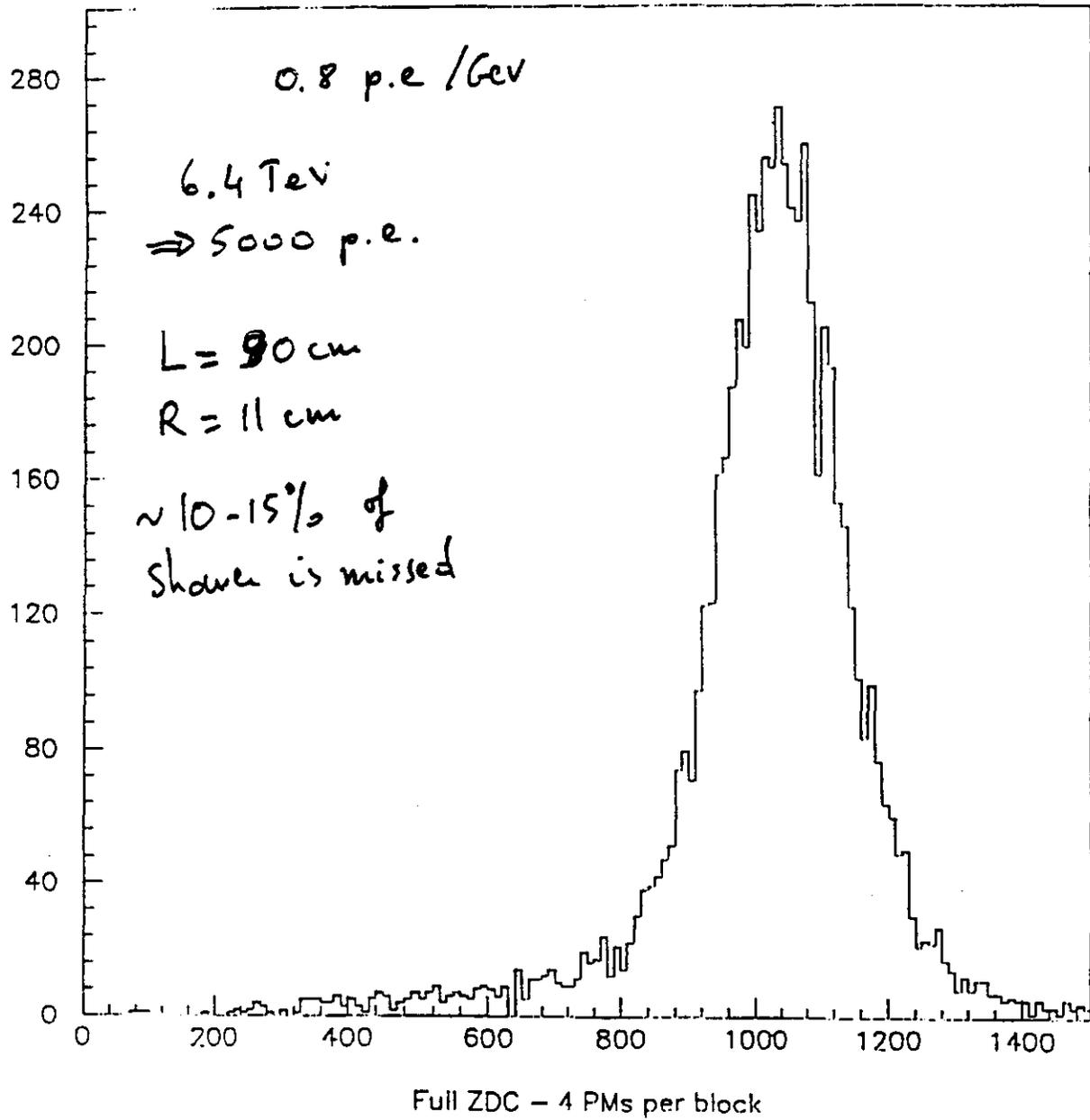


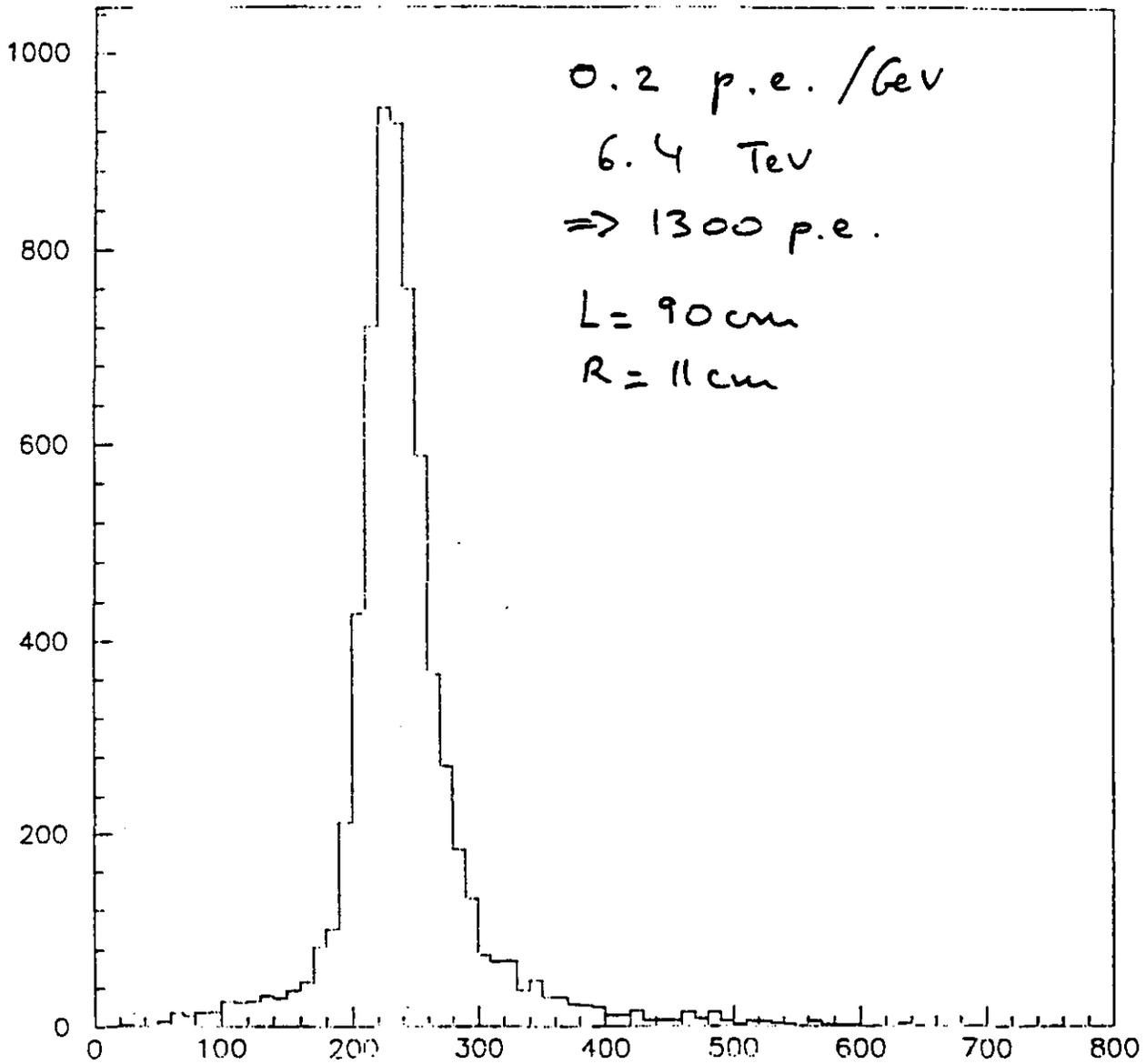
1.95 mm
 $(\frac{1}{32} \text{ "})$







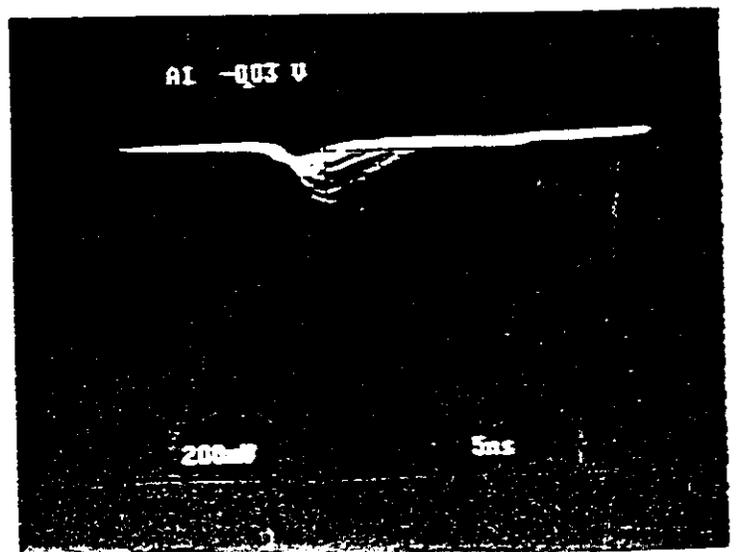
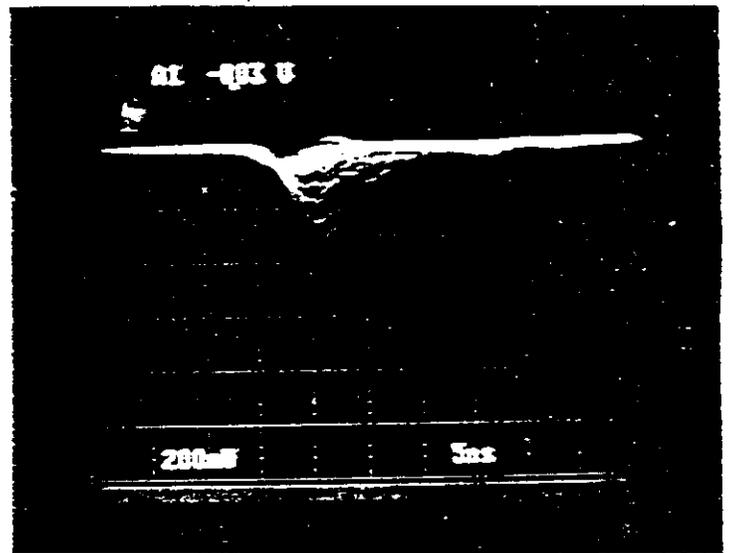
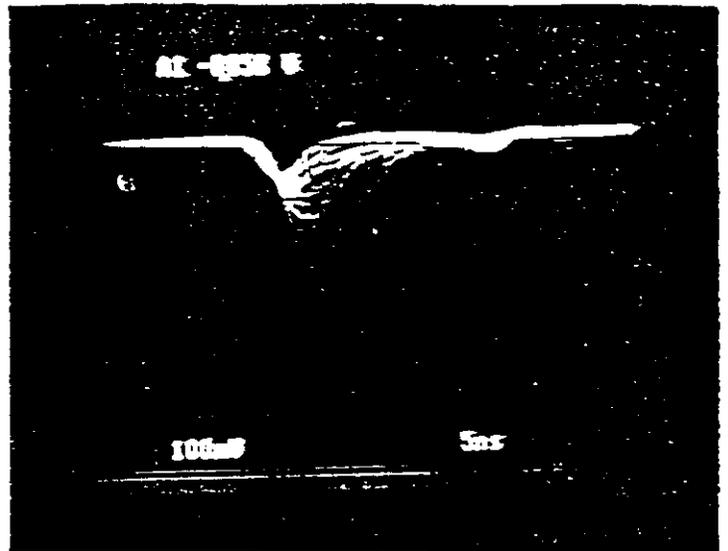




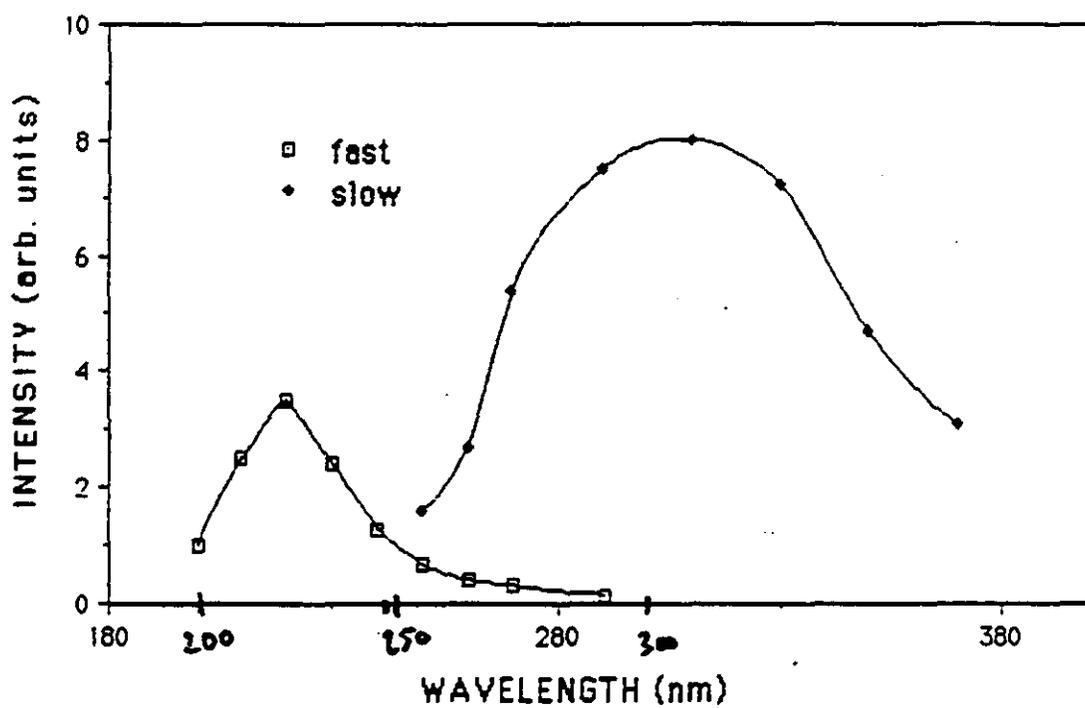
Full ZDC - 1 PM per block

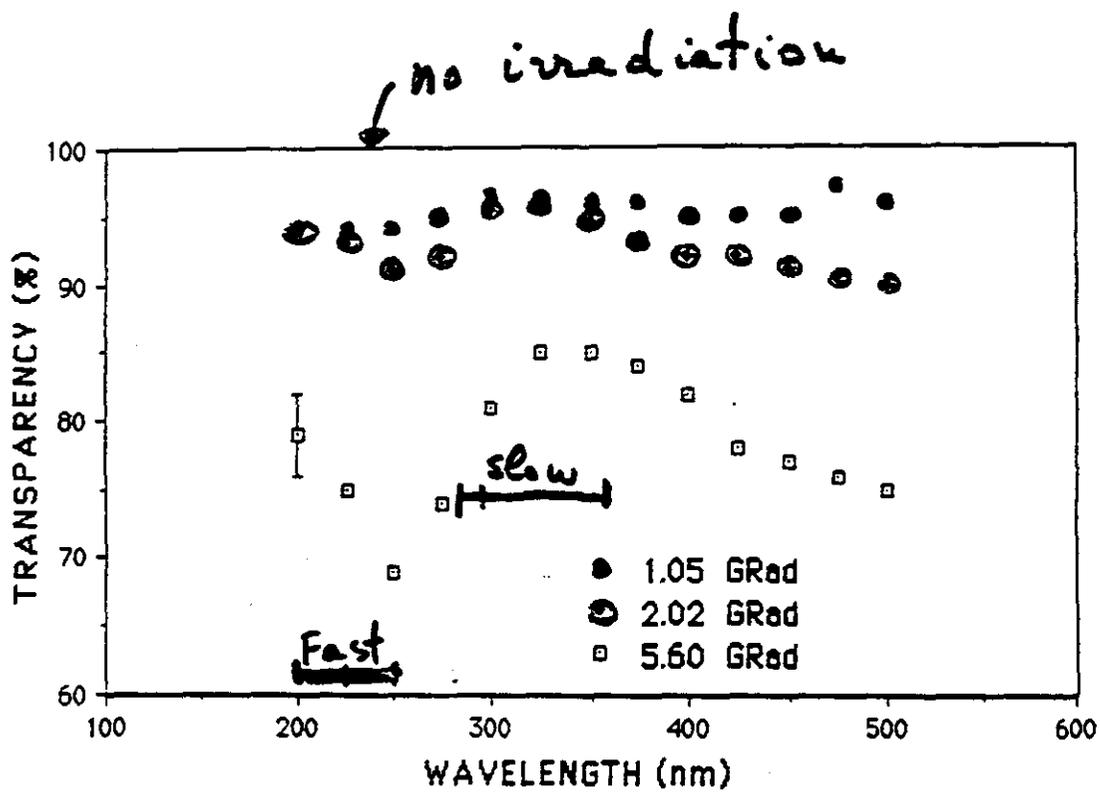
104/192.

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55V
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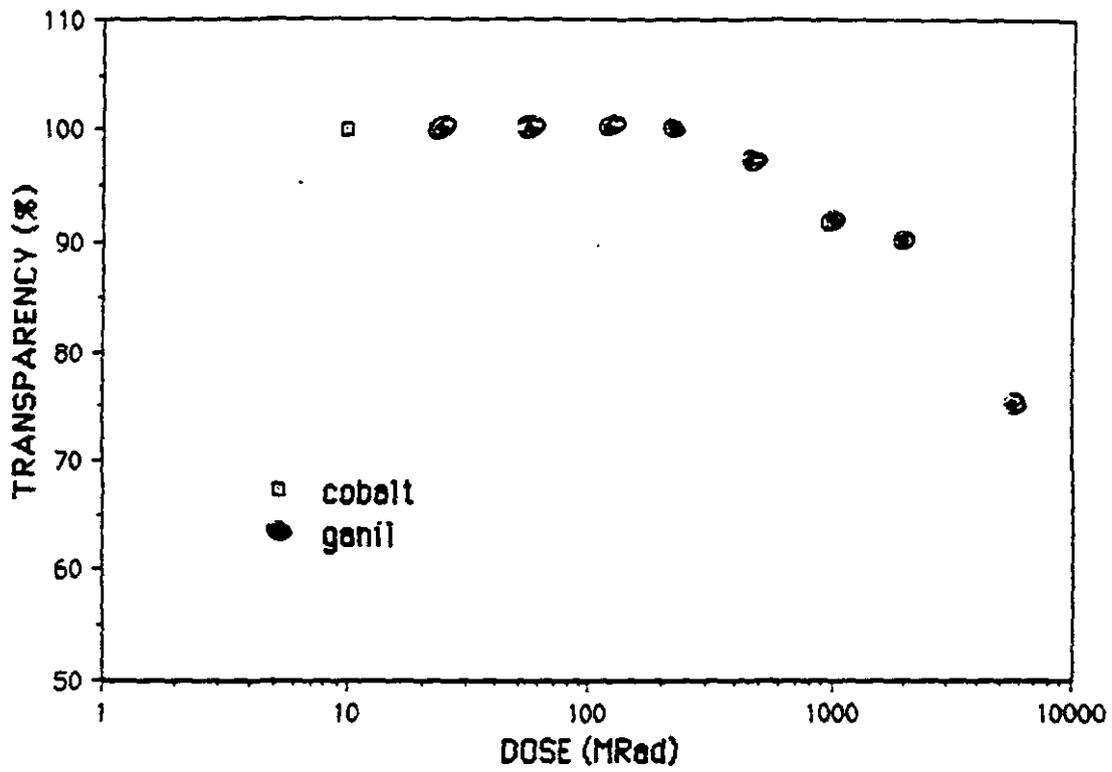


The 2 components of BaF_2

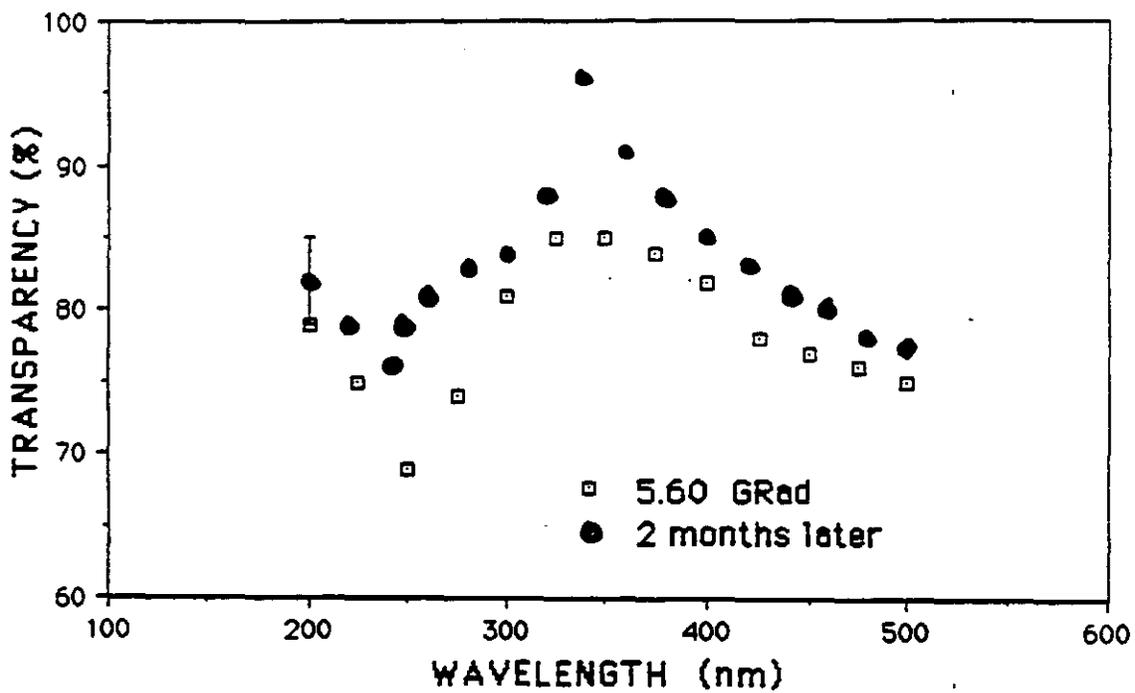




1 mm thick BaF₂



1 mm thick BaF_2
Fast component



1 mm thick BaF₂ .

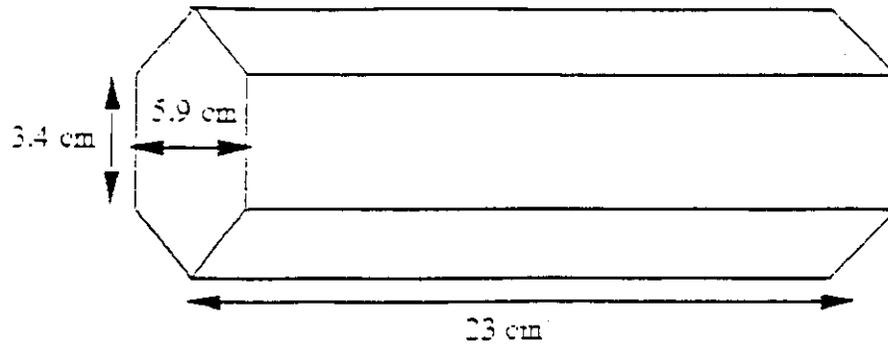


Fig V.7: Caractéristiques d'un module TAPS.

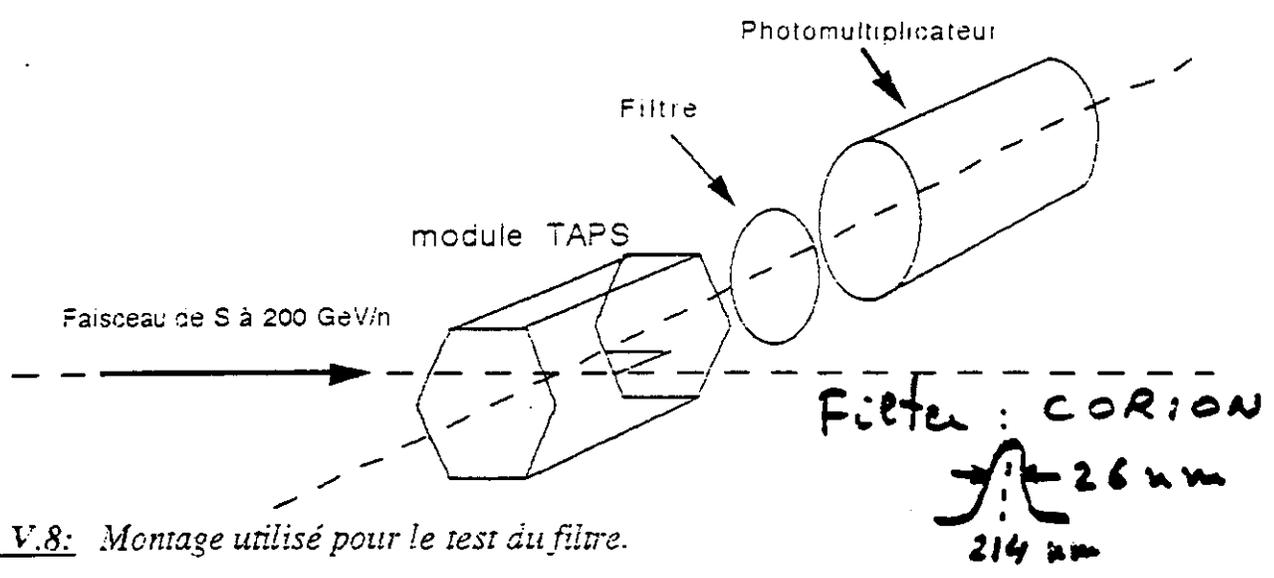


Fig V.8: Montage utilisé pour le test du filtre.

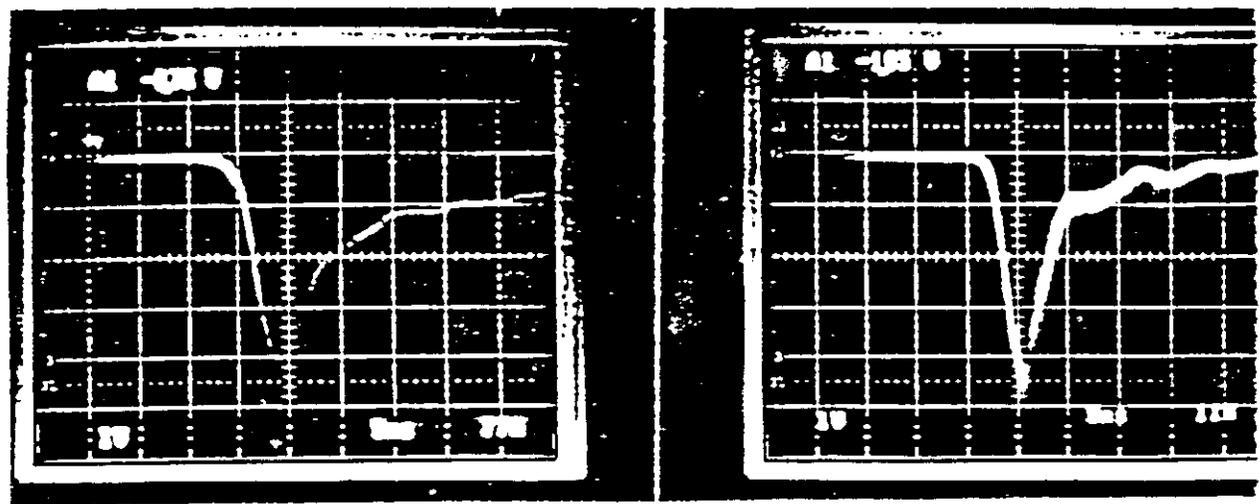
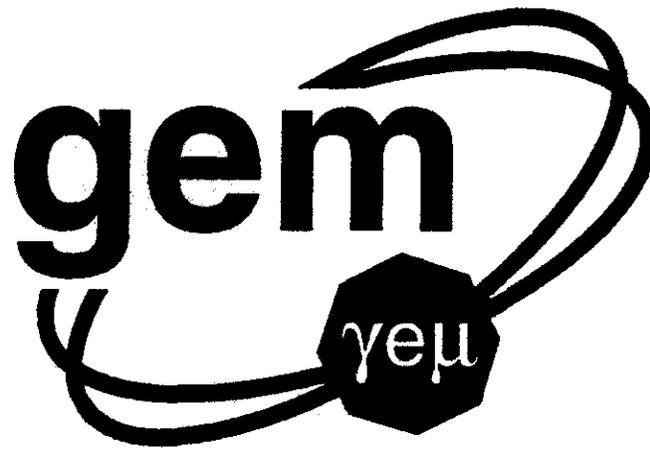


Fig V.9: Courbe de l'oscilloscope sans le filtre (à gauche) n°1 avec le filtre (à droite) n°2. Les échelles horizontale et verticale sont respectivement égales à 5ns par division et 1 V par division.

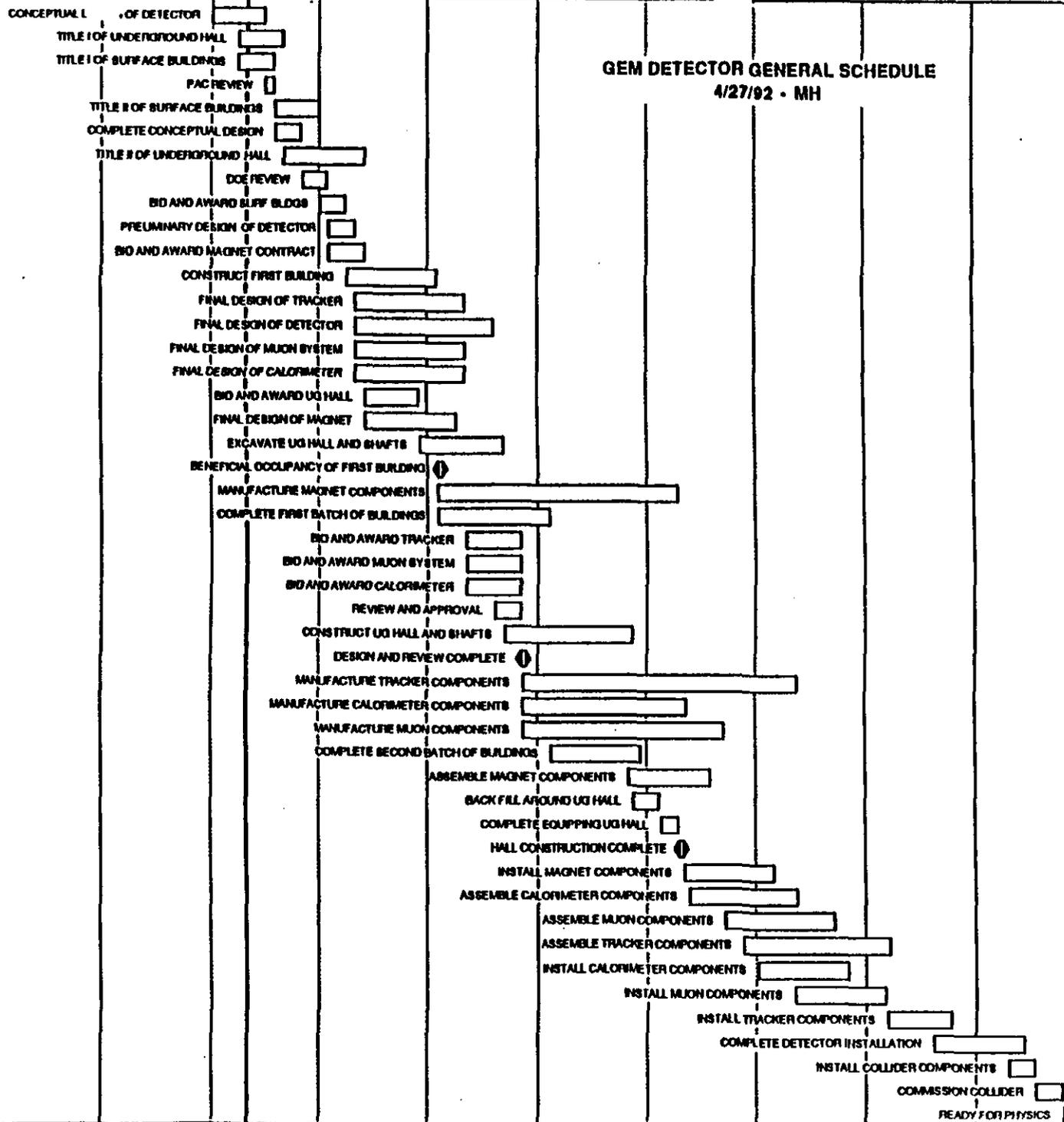
La composante lente apparaît sur la photographie n°1 à gauche du pic, le signal ne revient pas à 0 mais reste constamment à -1 V. Sur la photographie n°2, le signal revient à 0. Nous constatons bien l'effet du filtre. Cette atténuation du signal d'un



Presentation by:

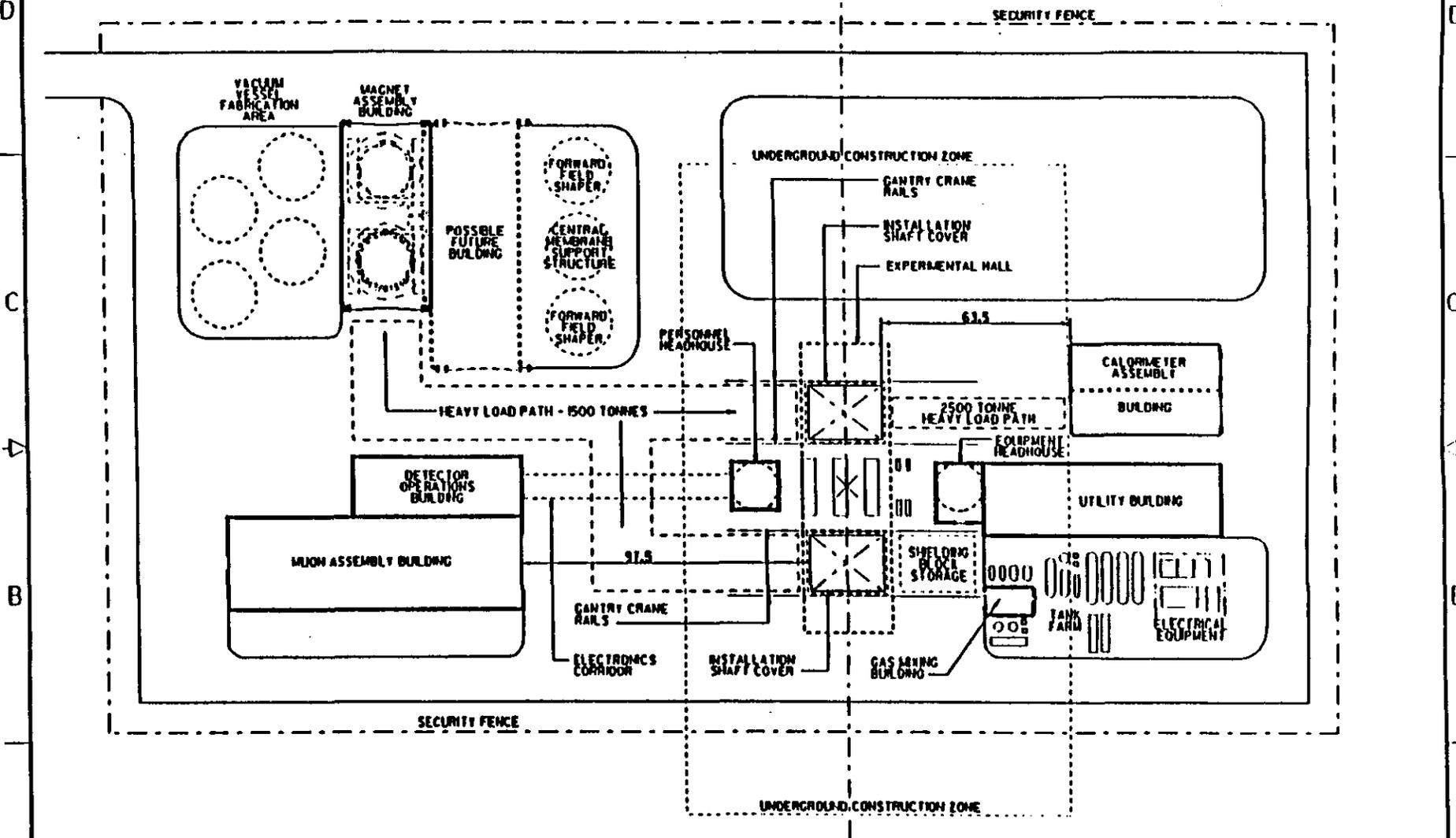
Mark Rennich

GEM DETECTOR GENERAL SCHEDULE 4/27/92 - MH





IR-6
↑



SITE PLAN

DIMENSIONS ARE SHOWN IN METERS UNLESS NOTED OTHERWISE.

THIS IS A CAD GENERATED DRAWING. DO NOT MAKE MANUAL REVISIONS OR ALTERATIONS.

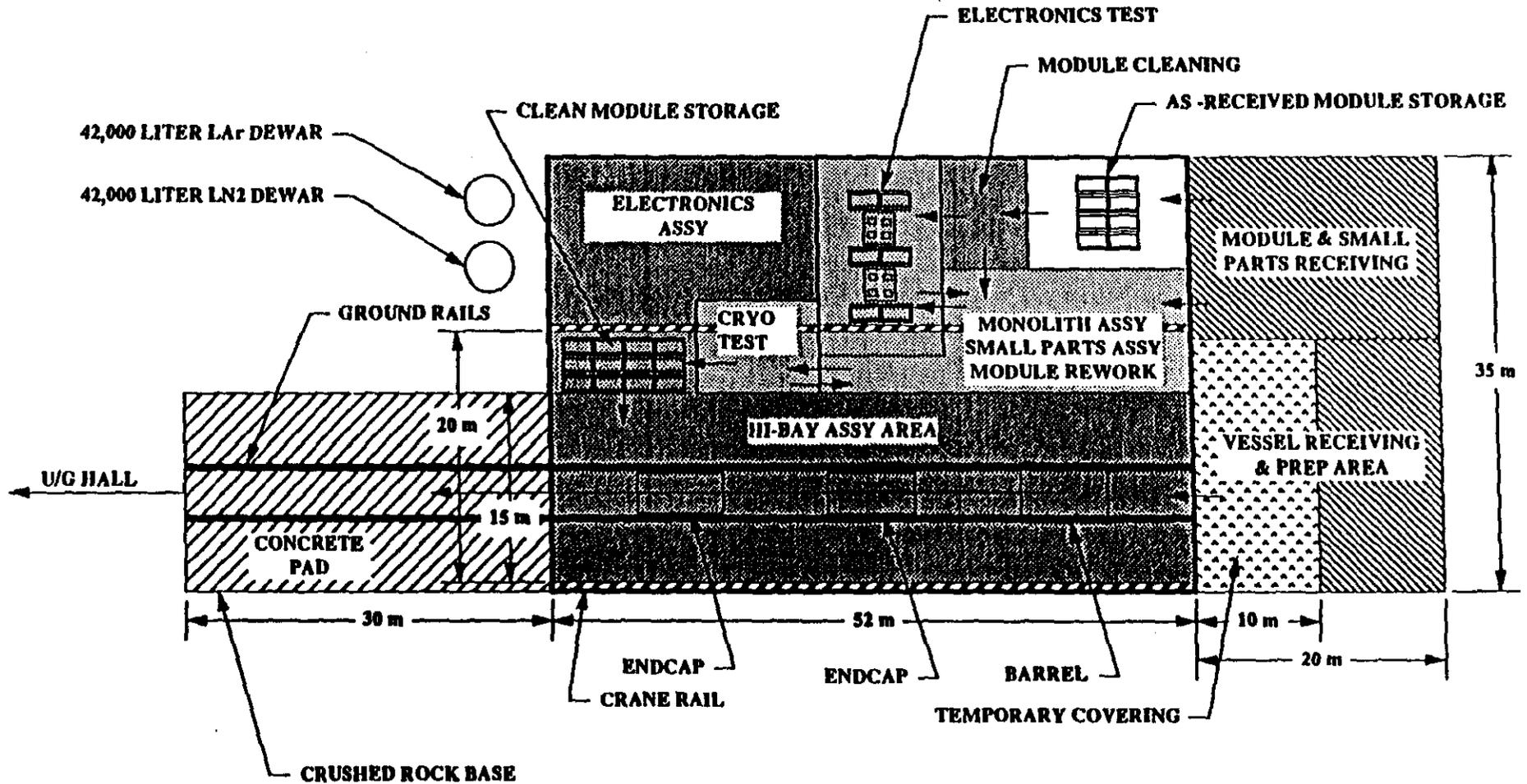
| | | | | | |
|-------------------------|--|--|-----------------------------|--|---------------------|
| | | | GEN EXPERIMENTAL FACILITIES | | |
| | | | IR-5 SITE PLAN | | |
| I. NAMES I. PROSAPID | | | CONTRACT NAME: N-K-2142-01 | | SHEET NO. 1 OF 1 |



PERMANENT BLDG BOUNDARY



CLEAN AREA LEGEND



CRANE REQUIREMENTS

HI-BAY: 50/10 TONNE BRIDGE CRANE
 VESSEL RECEIVING/PREP: 50 TONNE BOOM CRANE

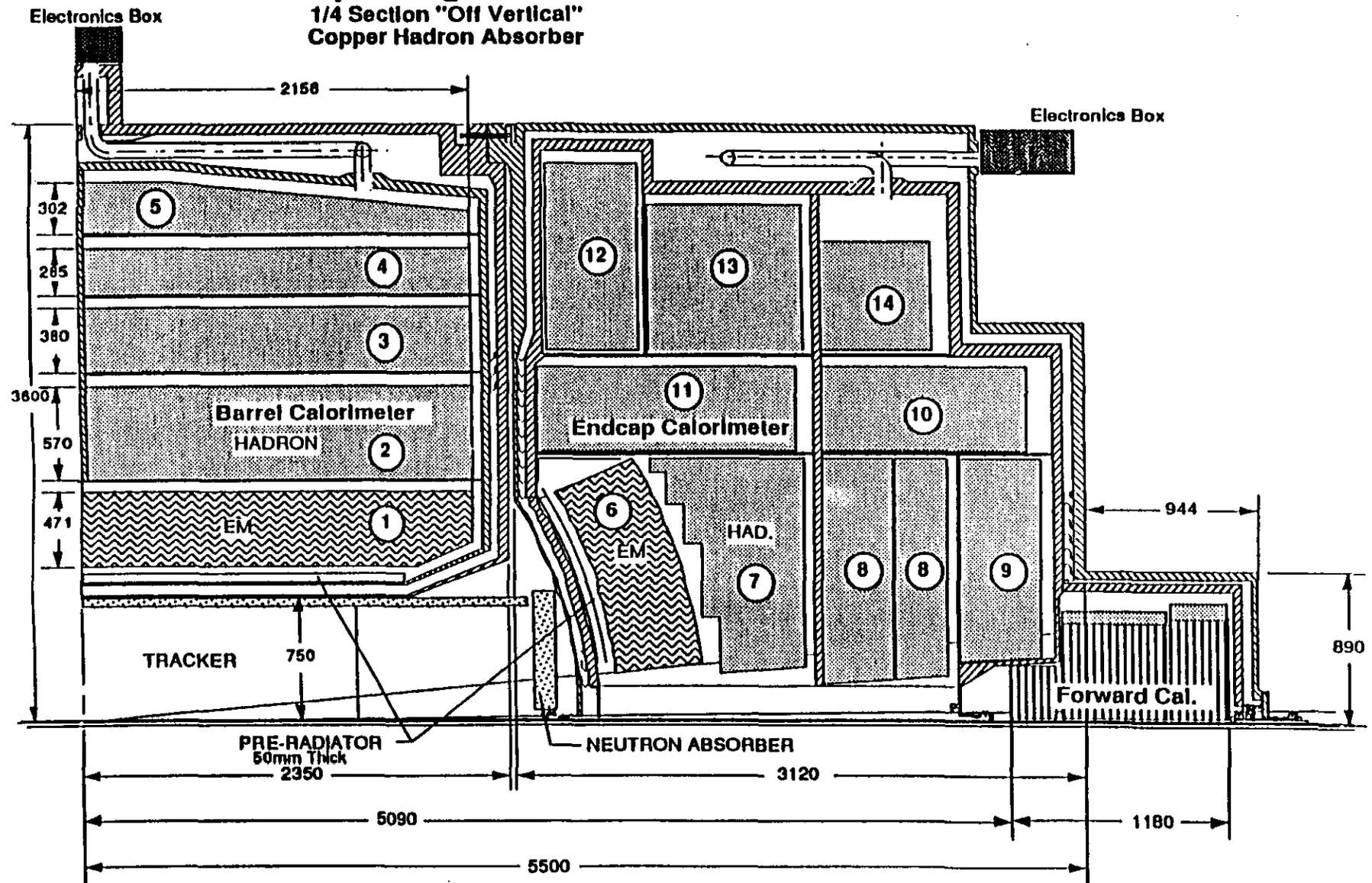
**GEM CALORIMETER ASSY BLDG
 PLAN VIEW**

DRAFT
 SCALE 500:1
 Eberle 03/02/92

Configuration

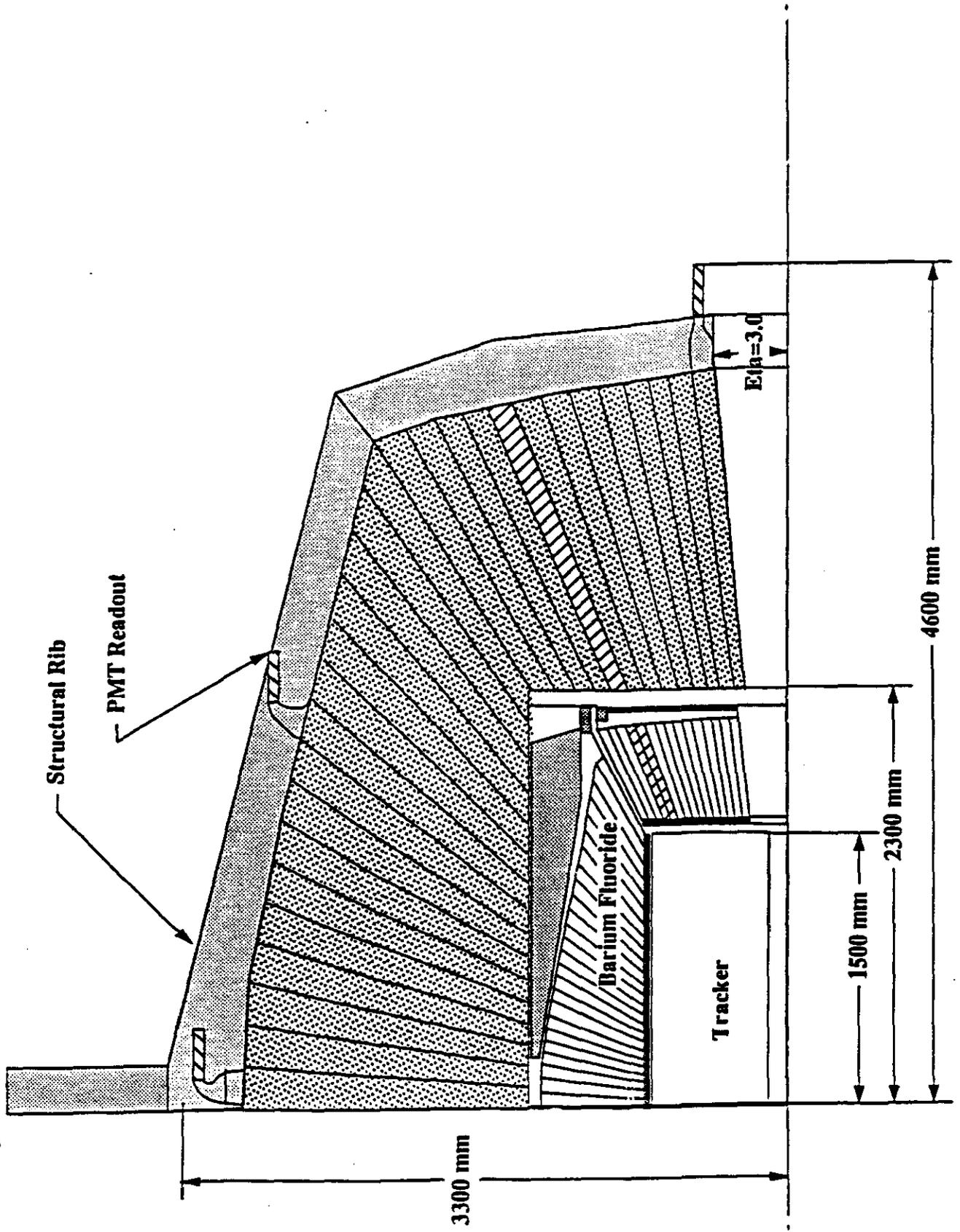
GEM Liquid Argon Calorimeter

1/4 Section "Off Vertical"
Copper Hadron Absorber

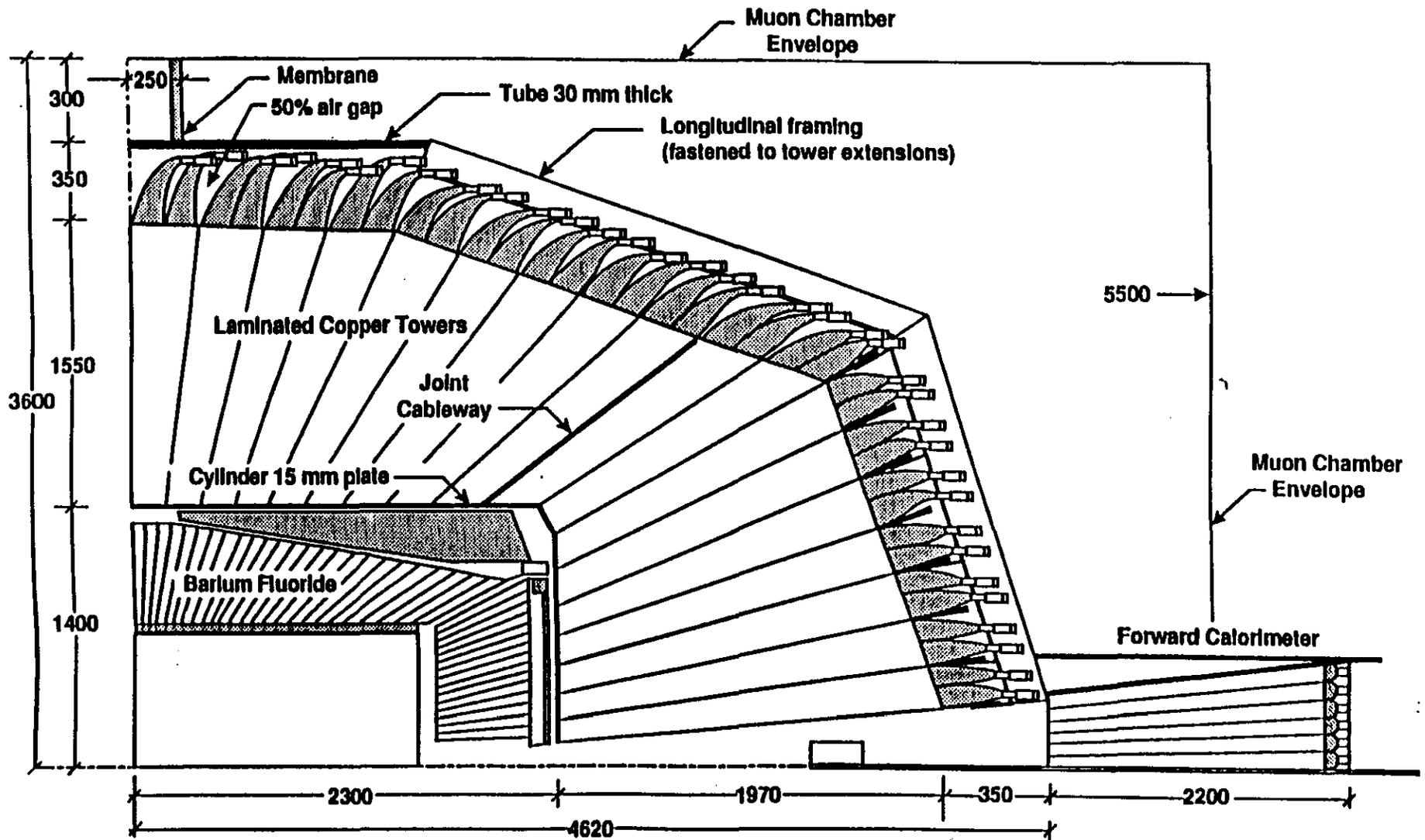


LLMason GEM LAC. 4/10/92

4/21/92

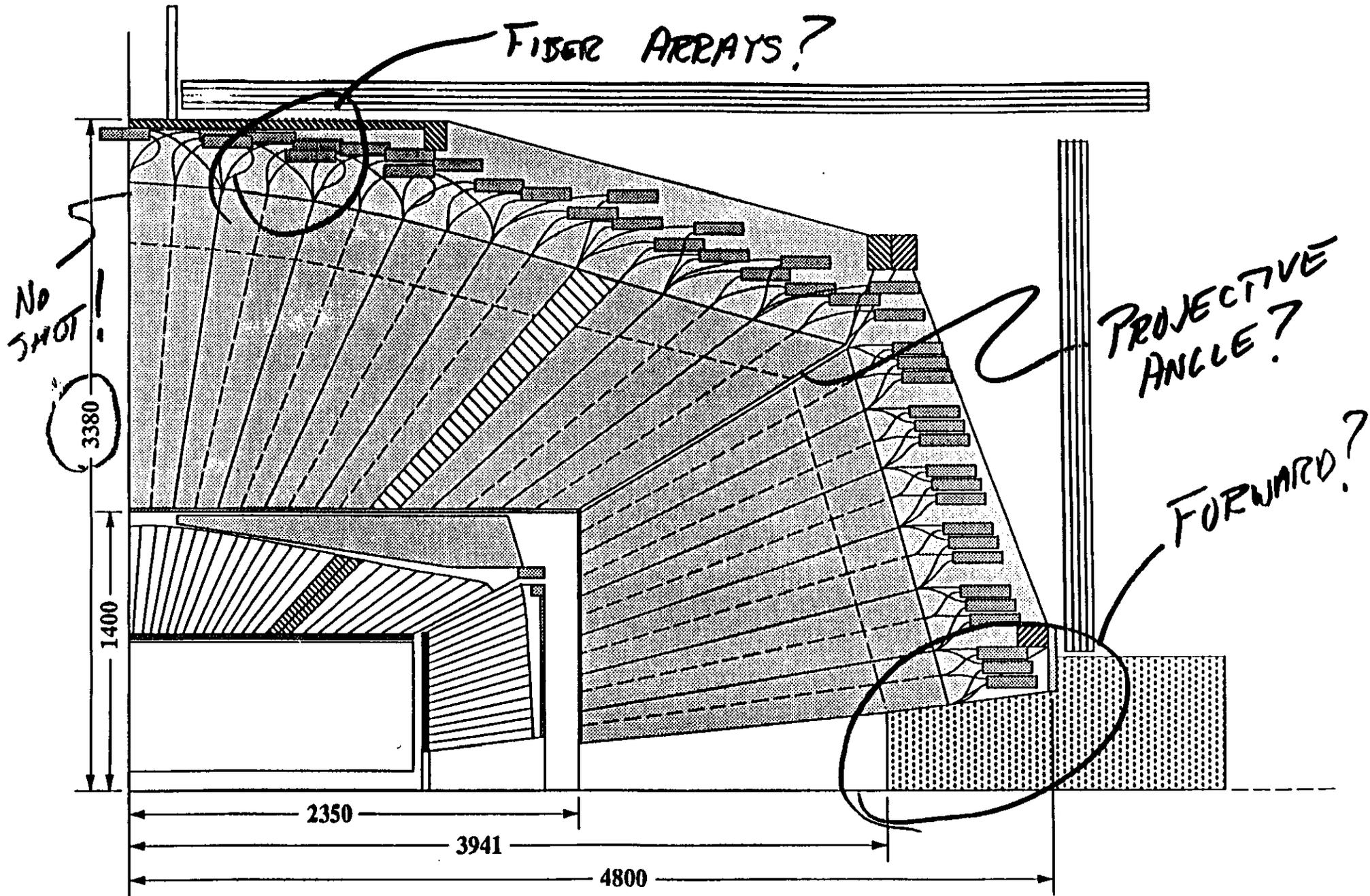


Spaghetti Hadron Calorimeter
 Copper Fill/0.12 Segmentation
 Barium Fluoride EM Calorimeter



Dimensions in millimeters

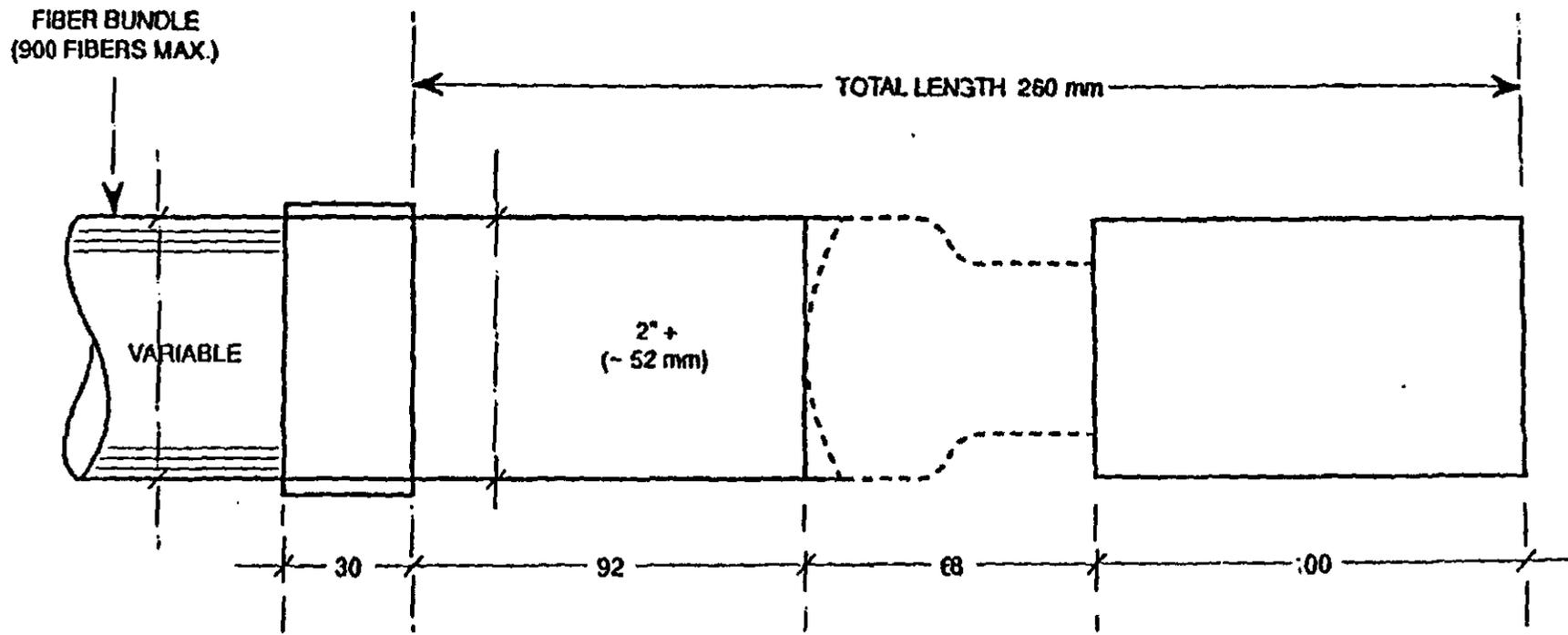
GEM Scintillator Calorimeter System



GEM Detector
Fiber Hadron Calorimeter

LIGHT GUIDE / PMT / BASE ASSEMBLY

SCHEMATIC VIEW



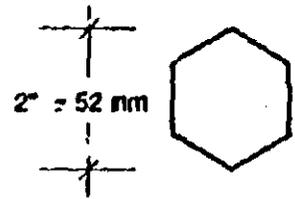
CLAMP
HEX

LIGHT MIXER
HEXAGONAL

PMT
2" TUBE #2490

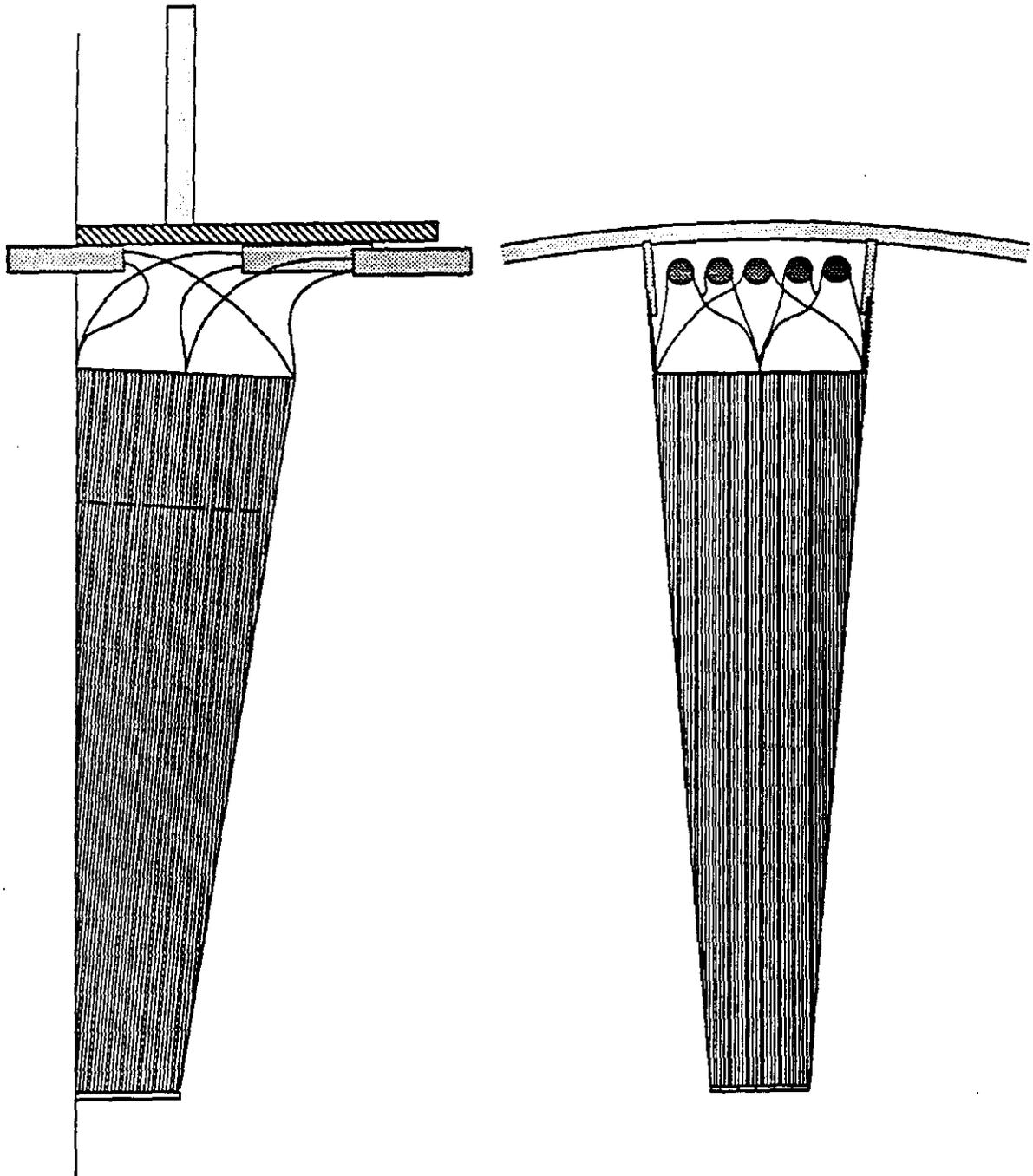
PMT BASE

2" APEX - APEX



L = 38 mm
φ = 51 mm

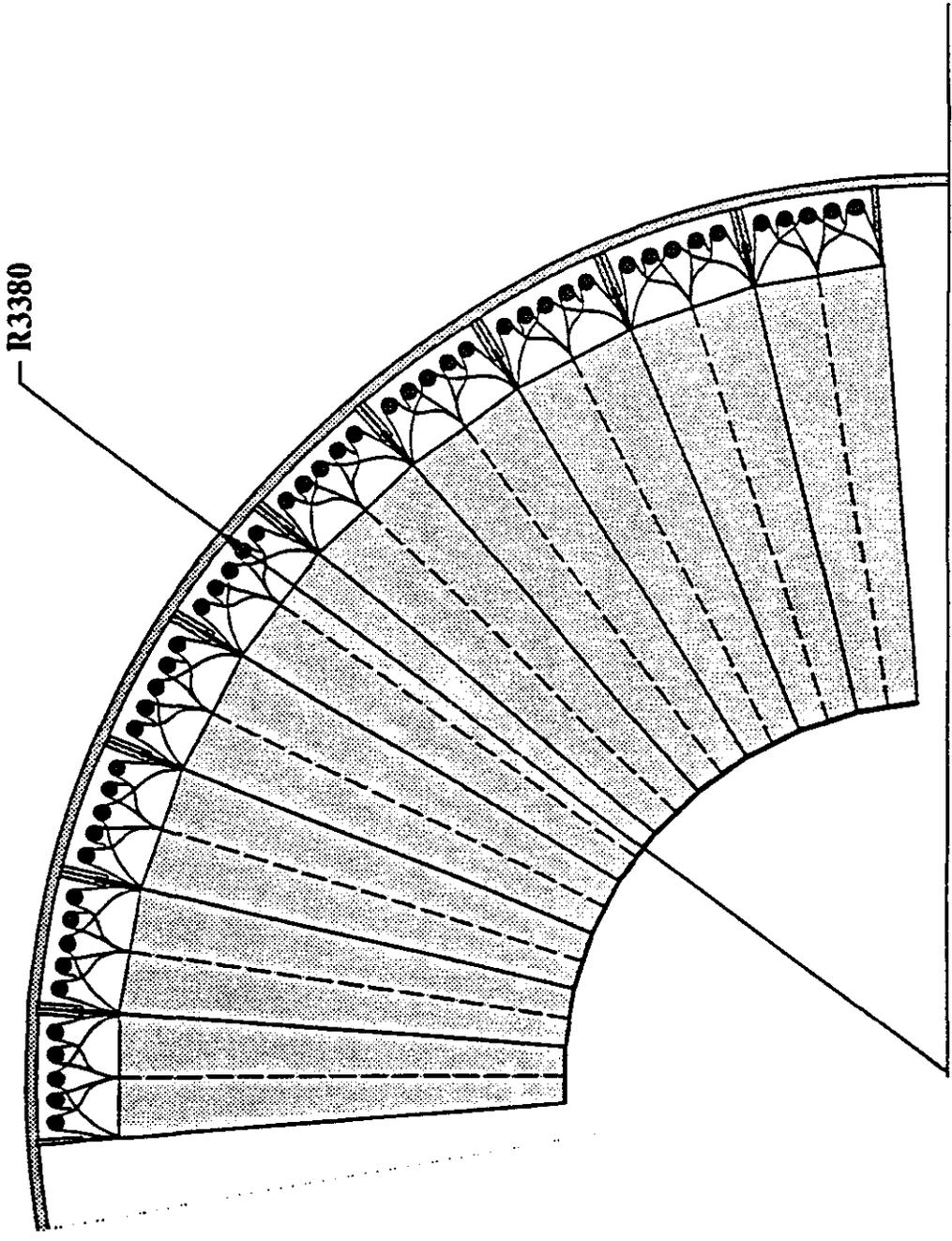
DIMENSIONS IN MILLIMETERS



**GEM Detector
Fiber Hadron Calorimeter
Eta=0 Tower**

G.03.CU.021

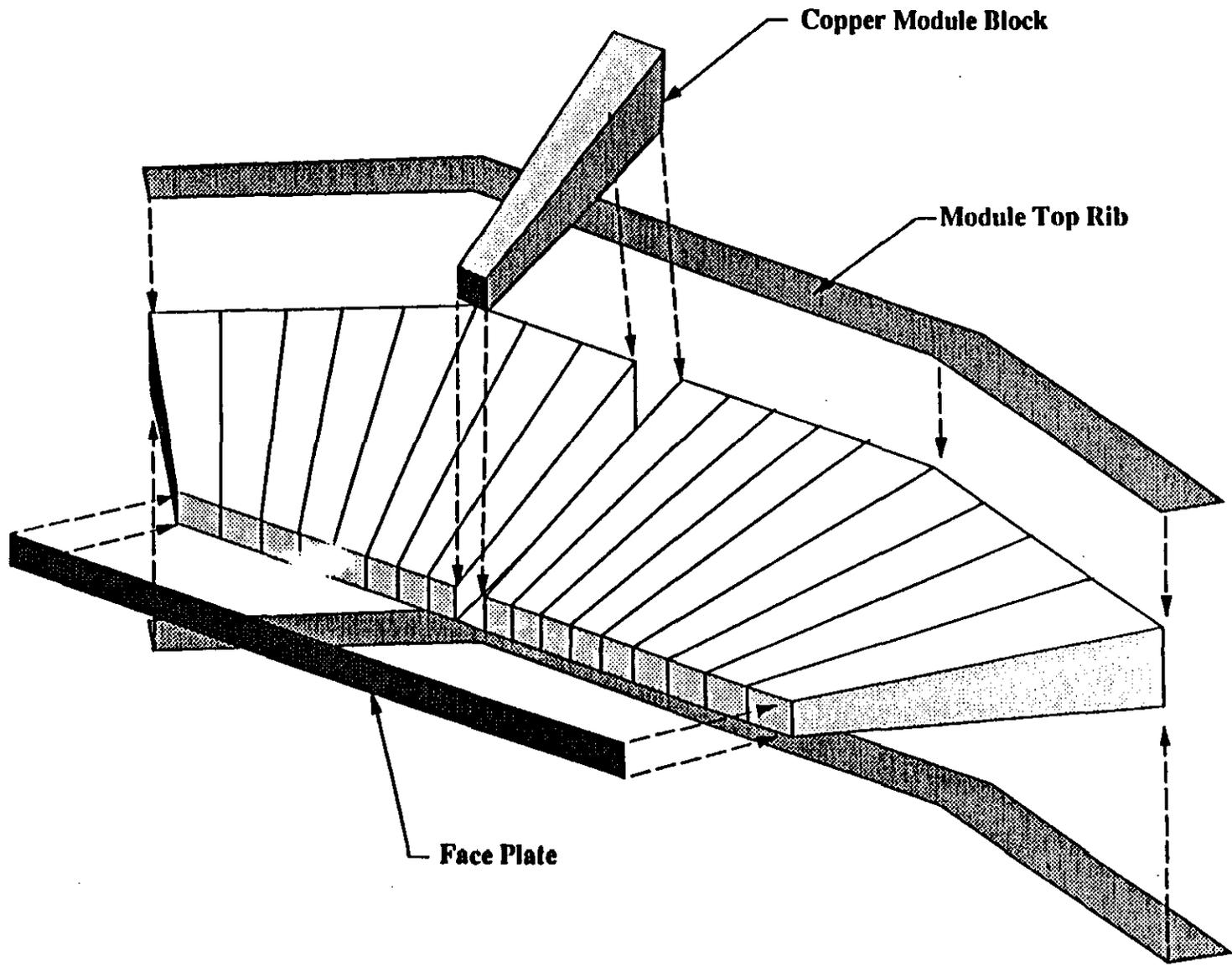
Rennich



**GEM Detector
Fiber Hadron Calorimeter**

Computed Parameters for Complete Fiber Calorimeter

| Total | | |
|---------------------------|-----------|----------------|
| Number of Mech.Theta Rows | 28 | Each |
| Number of Mech. Towers | 940 | Each |
| Absorber Volume | 169.88 | M ³ |
| Fiber Length | 3,749,264 | M |
| Cu Tube Length | 2,884,050 | M |
| Cu Tube Volume | 131.00 | M ³ |
| Number of Cu Plates | 48,668 | each |
| Copper Plate Weight | 1292.47 | Mt |
| Number of Fibers | 2,510,848 | each |
| Total Weight | 1434.99 | MT |



**GEM Detector
Fiber Calorimeter
Central Barrel Module
Assembly-Station One**

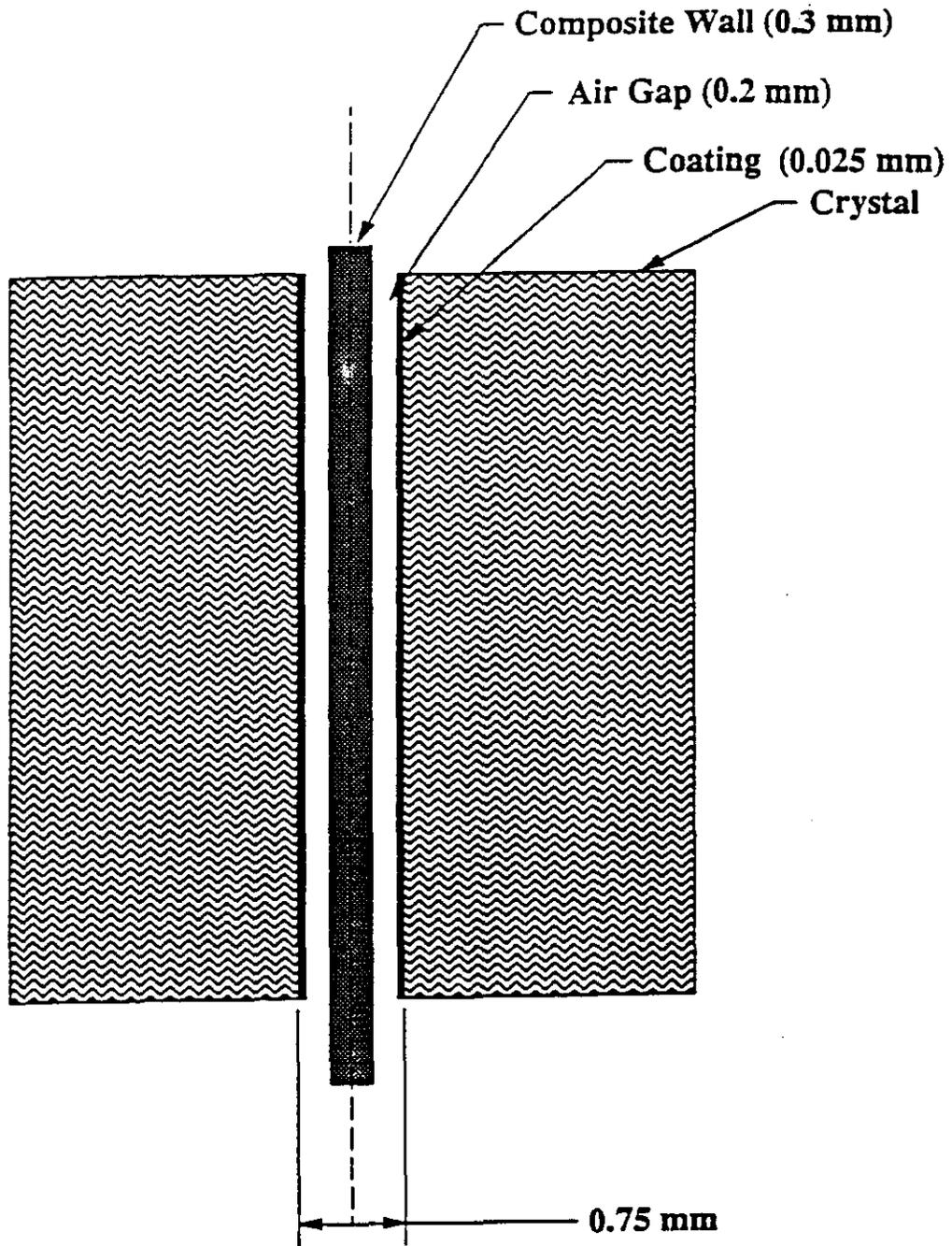
Scintillating Fiber Calorimeter
Category Distribution

| Category | Estimate | Fraction |
|------------------------------|------------------|-----------------|
| Research & Devel. | \$3,445K | 11.3% |
| EDIA | \$9,419K | 30.9% |
| Construction | \$30,492K | |
| Contingency | \$12,375K | 31.0% |
| Total | \$55,730K | |

GEM Barium Fluoride EM Calorimeter

Potential Improvements

| | Current | Potential |
|-----------------------|-----------------------------|-------------------------------------|
| Structure | Fiber Composite | Titanium/Aluminide |
| Tolerancing | Centerline Reference | Face Reference |
| Inspection | Mechanical Gages | Coordinate Measuring Machine |
| Readout | Phototriode | Solid State Diode |
| Joint Matching | Lapping | Diamond Cutting |

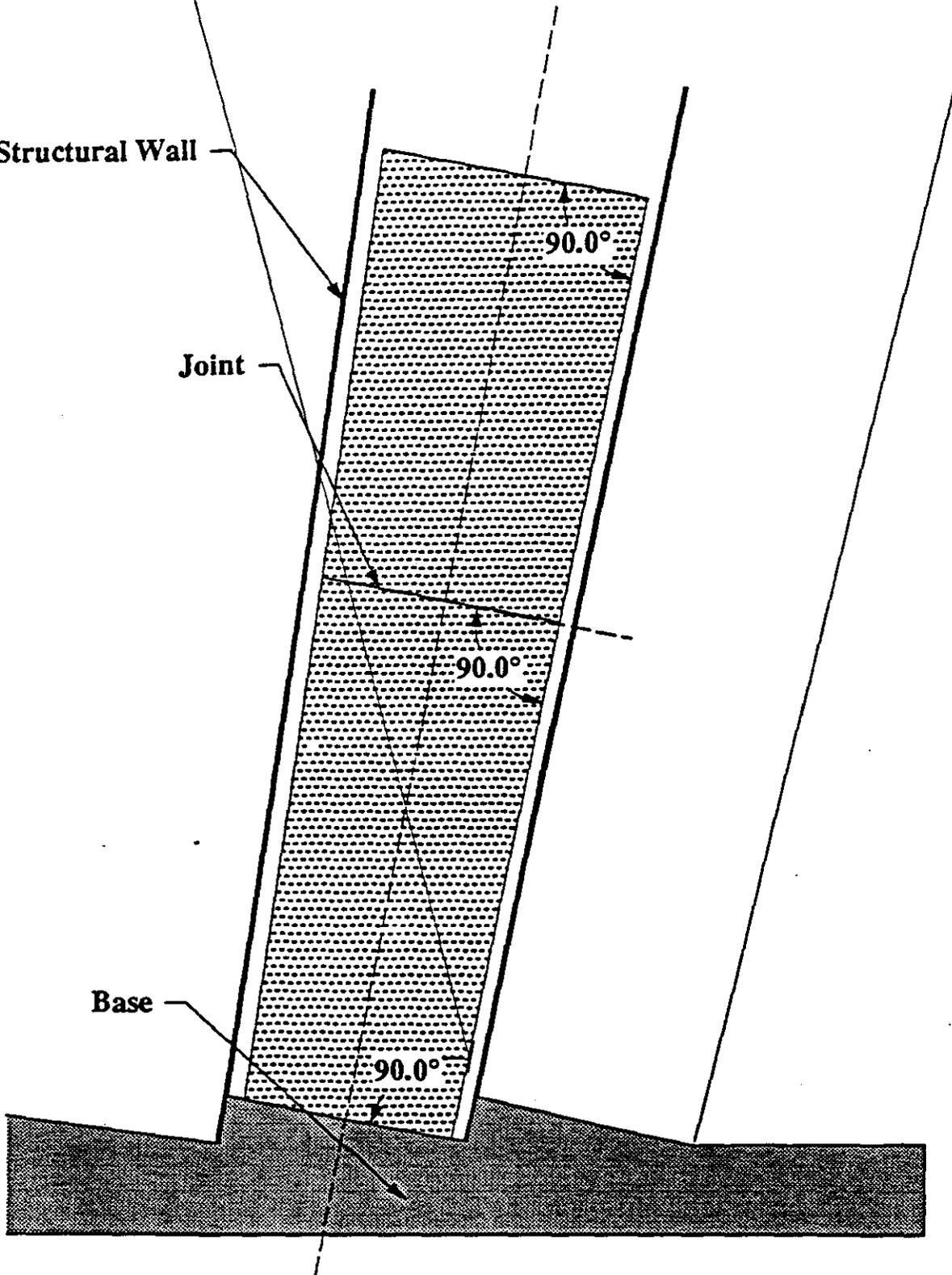


**Barium Fluoride
Detail of Gap Between Crystals**

Structural Wall

Joint

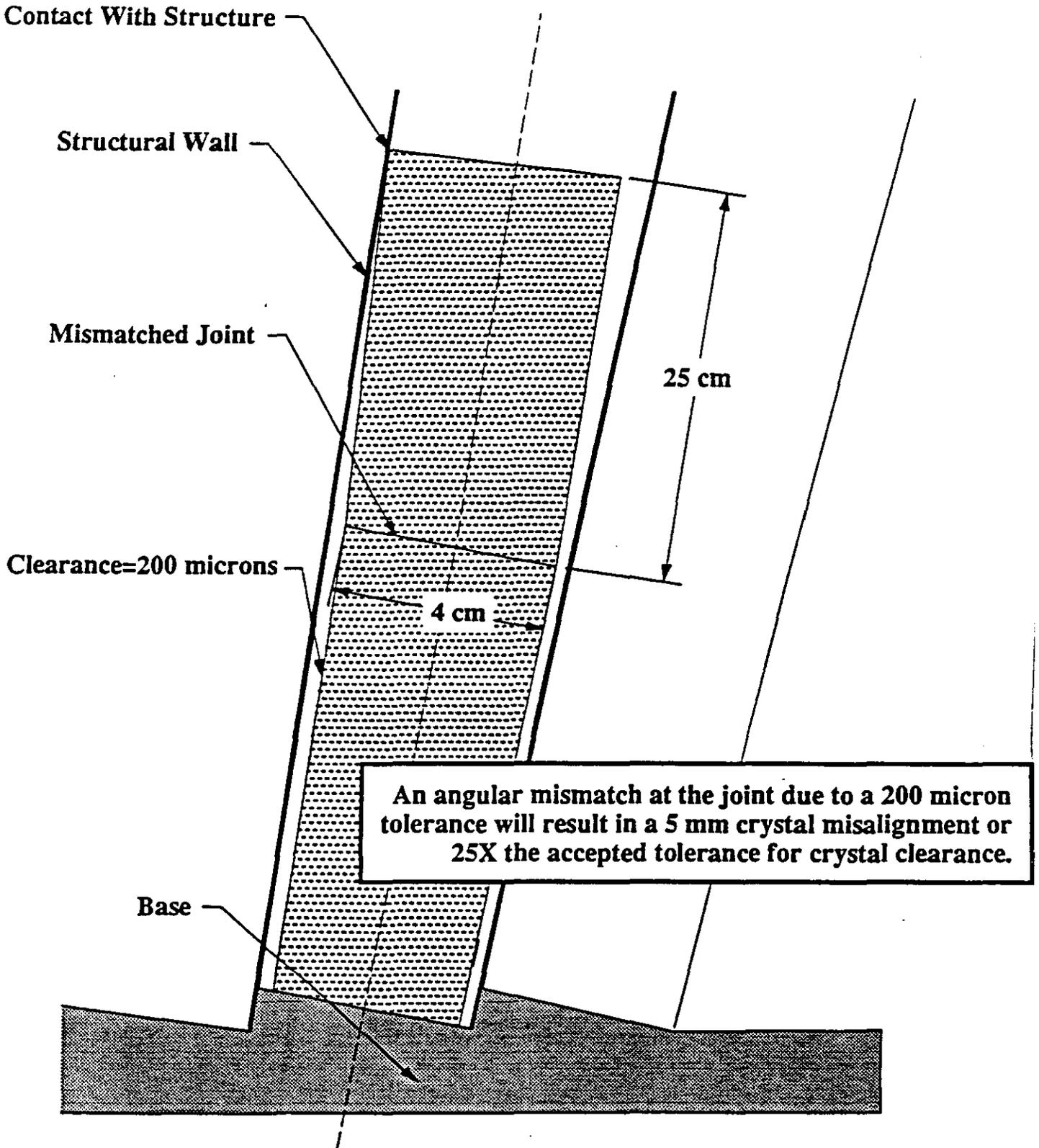
Base



**Barium Fluoride
Preferred Crystal Tolerancing**

G.03.BF.010

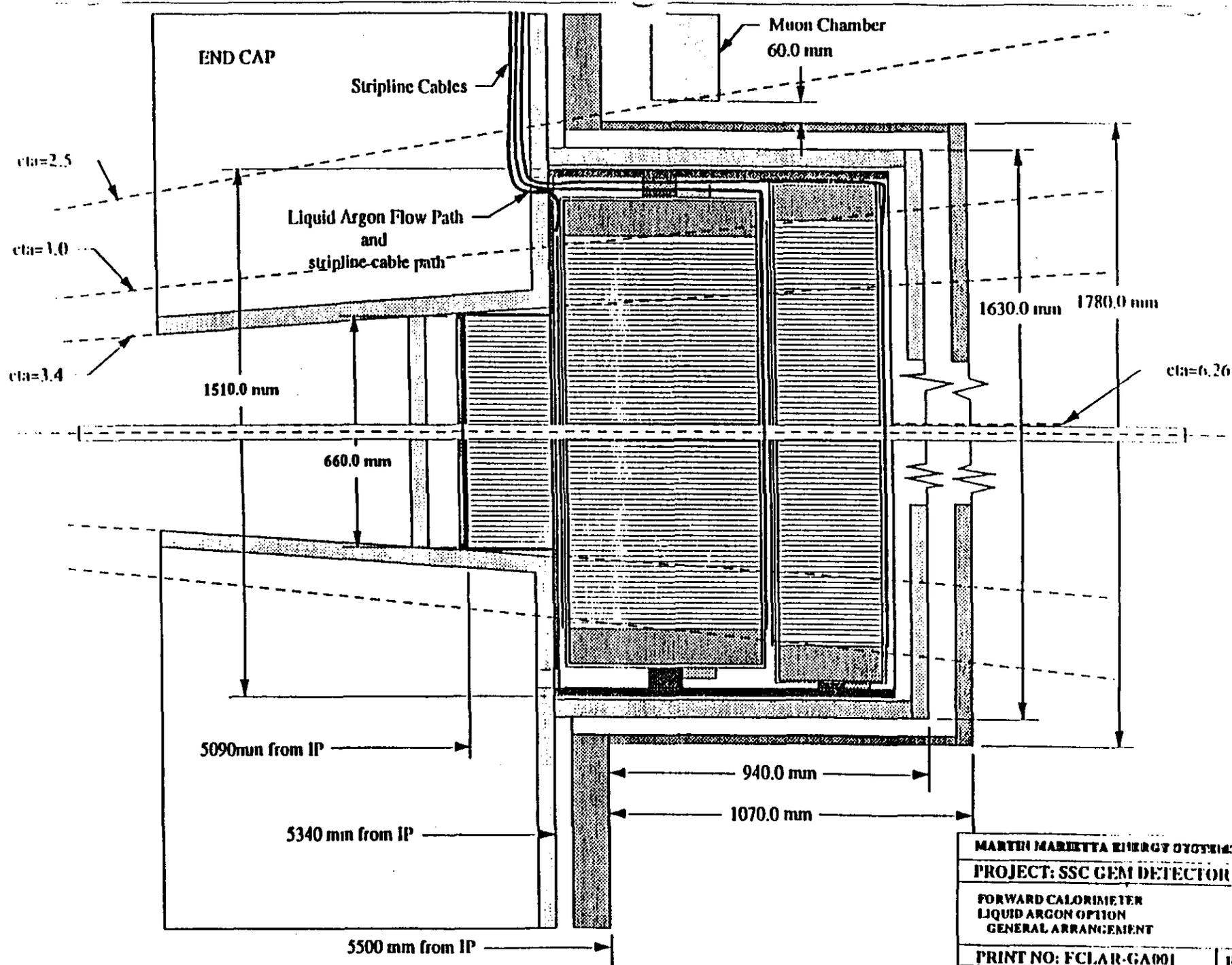
Rennich



**Barium Fluoride
 Crystal Tolerancing
 Joint Error**

Barium Fluoride Calorimeter
Category Distribution

| Category | Estimate | Fraction |
|------------------------------|------------------|-----------------|
| Research & Devel. | \$4,093K | 7.2% |
| EDIA | \$8,271K | 14.6% |
| Construction | \$56,509K | |
| Contingency | \$11,417K | 17.6% |
| Total | \$80,290K | |



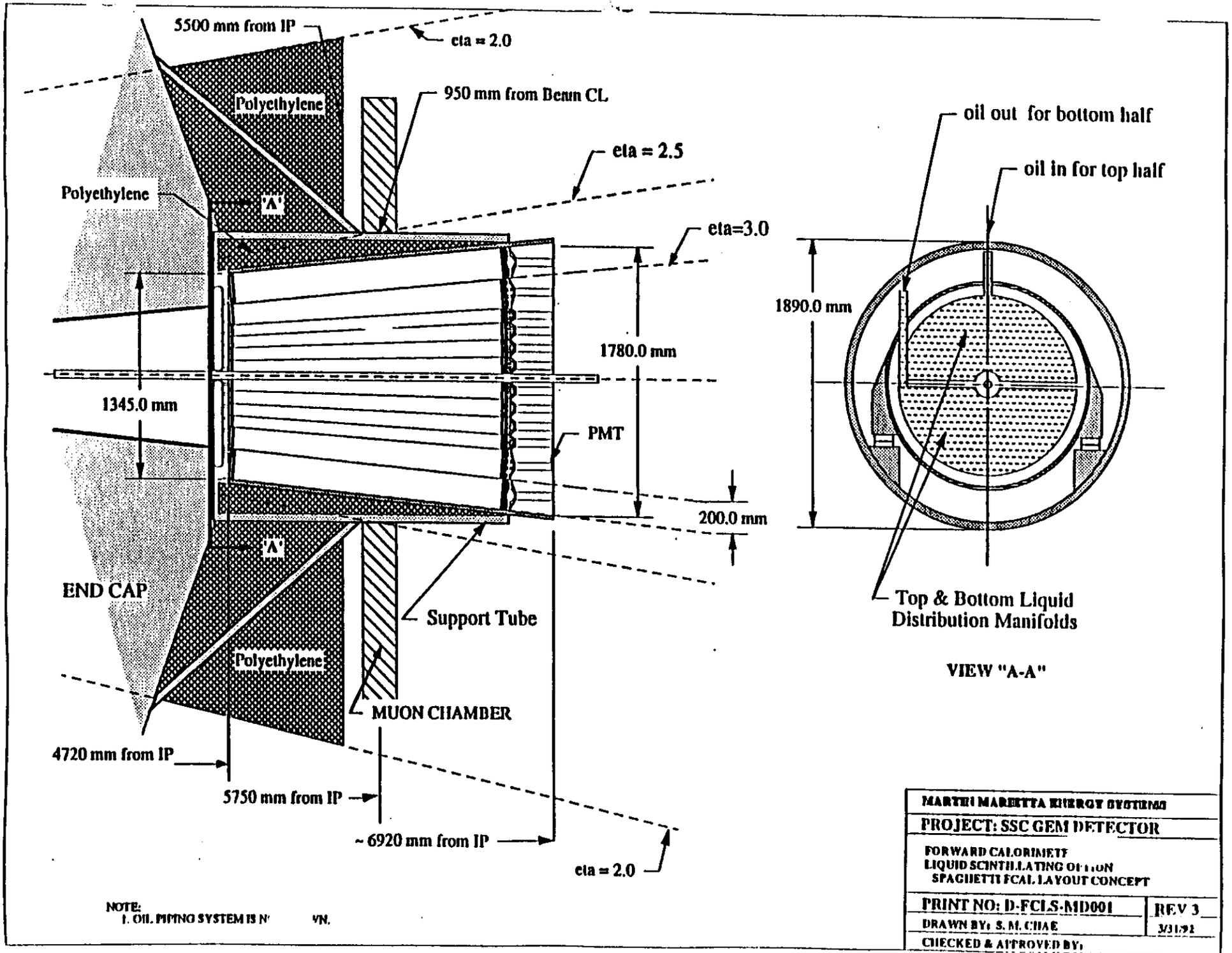
| | |
|---|-------|
| MARTIN MARIETTA ENERGY SYSTEMS | |
| PROJECT: SSC GEM DETECTOR | |
| FORWARD CALORIMETER LIQUID ARGON OPTION GENERAL ARRANGEMENT | |
| PRINT NO: FCLAR-GA001 | REV 4 |
| DRAWN BY: S. M. CHAF | 4/01 |
| CHECKED & APPROVED BY: | |

LAR COST SUMMARY

LIQUID ARGON OPTION FORWARD CALORIMETER
COST ESTIMATE SUMMARY

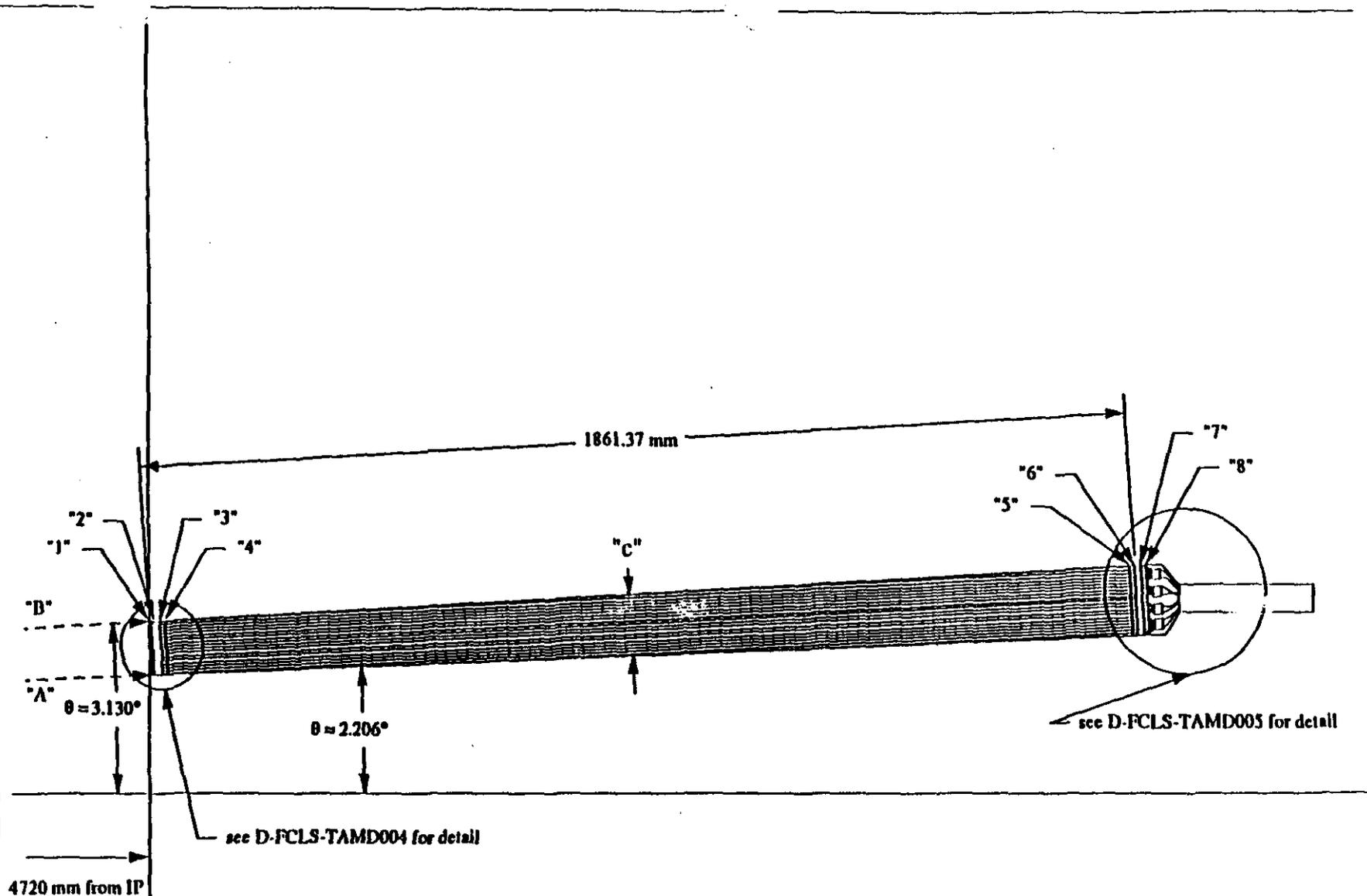
| | ENGR/DESIGN M&S and INSPECTION | PROC/FAB MATERIAL | LABOR ASSEMBLY | LABOR INSTALLATION | SUBTOTAL |
|--------------|--------------------------------------|----------------------|-------------------|-----------------------|-----------|
| CONSTRUCTION | \$2,332K | \$7,103K | \$883K | \$20K | \$10,338K |
| R&D | \$708K | \$165K | \$37K | \$0K | \$910K |
| ENGINEERING | \$282K | \$0K | \$0K | \$0K | \$282K |
| SUBTOTAL | \$3,323K | \$7,268K | \$920K | \$20K | \$11,530K |

| | COST | PERCENTAGE |
|--------------|-----------|------------|
| R&D | \$910K | 11.37% |
| EDIA | \$2,614K | 32.66% |
| CONSTRUCTION | \$8,006K | |
| CONTINGENCY | \$3,756K | 35.37% |
| TOTAL | \$15,286K | |



NOTE:
1. OIL PIPING SYSTEM IS N° VN.

| | |
|--|---------|
| MARTI MARETTA ENERGY SYSTEMS | |
| PROJECT: SSC GEM DETECTOR | |
| FORWARD CALORIMET LIQUID SCINTILLATING OIL SPAGHETTI CAL. LAYOUT CONCEPT | |
| PRINT NO: D-FCLS-MD001 | REV 3 |
| DRAWN BY: S. M. CHAE | 3/31/92 |
| CHECKED & APPROVED BY: | |



| | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |
|------------|---------|---------|---------|---------|---------|---------|---------|---------|
| RA | 181.90 | 182.09 | 182.67 | 183.06 | 253.17 | 253.55 | 253.59 | 253.98 |
| RB | 258.03 | 258.30 | 259.12 | 259.67 | 359.13 | 359.67 | 359.73 | 360.28 |
| **Length A | 4725.28 | 4730.28 | 4745.28 | 4755.28 | 6576.65 | 6586.65 | 6587.65 | 6597.65 |
| **Length B | 4725.28 | 4730.28 | 4745.28 | 4755.28 | 6576.65 | 6586.65 | 6587.65 | 6597.65 |
| Length C | 74.75 | 74.83 | 75.07 | 75.22 | 104.04 | 104.19 | 104.21 | 104.37 |

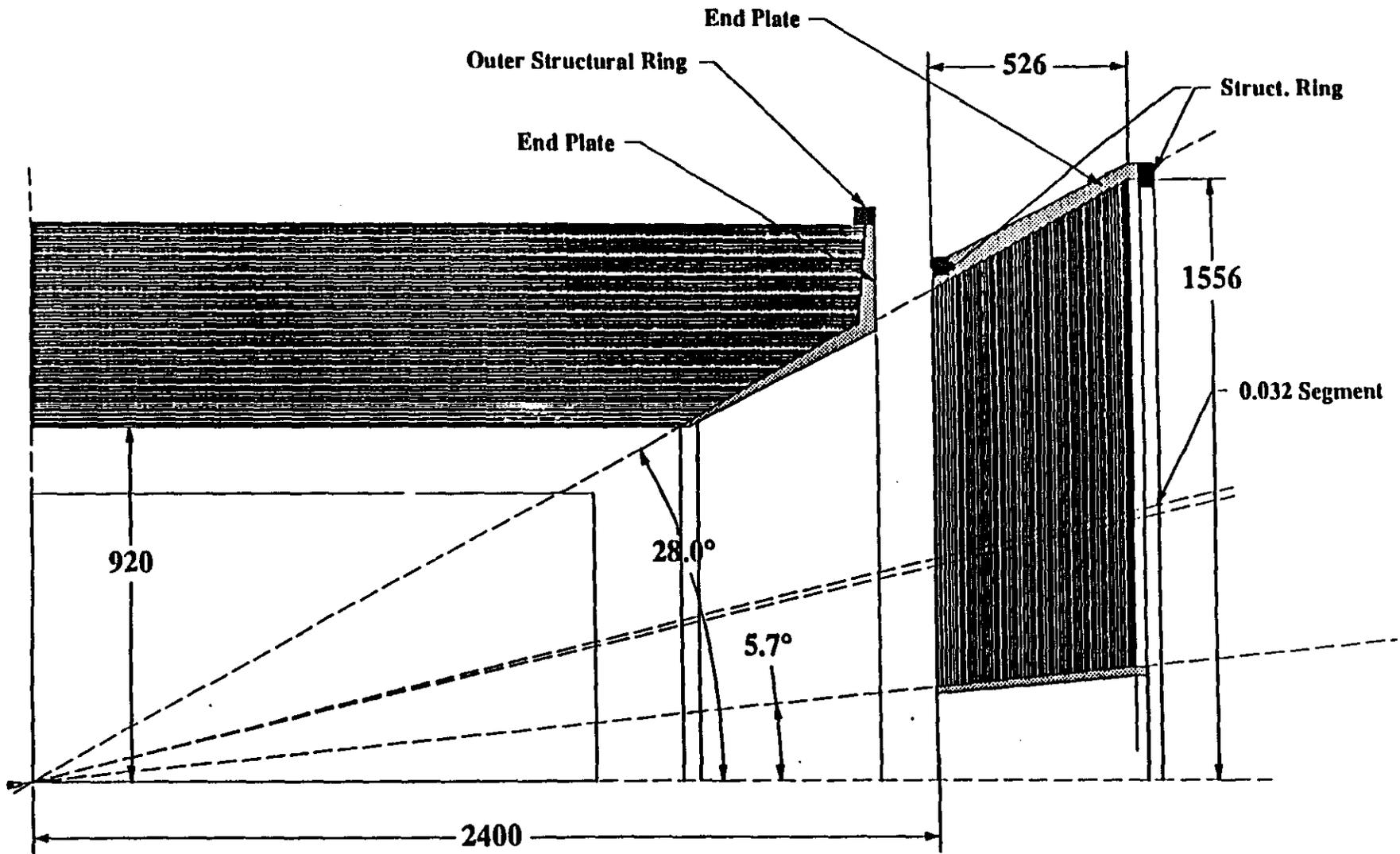
** The lengths in the table indicate the distance from the interaction point.

| | |
|--|--------|
| MARTINI MARETTA ENERGY SYSTEMS | |
| PROJECT: SSC GEM DETECTOR | |
| FORWARD CALORIMETER LIQUID SCINTILLATING OPTION SPAGHETTI TOWER & MODULE DESIGN SUPER TOWER OF MODULE 4 | |
| PRINT NO: D-FCLS-TAMD003 | REV 2 |
| DRAWN BY: S. M. CHAE | 3/2/93 |
| CHECKED & APPROVED BY: | |

SCINTILLATING LIQUID OPTION FORWARD CALORIMETER
COST ESTIMATE SUMMARY

| | ENGR/DESIGN M&S and INSPECTION | PROC/FAB MATERIAL | LABOR ASSEMBLY | LABOR INSTALLATION | SUBTOTAL |
|--------------|--------------------------------------|----------------------|-------------------|-----------------------|-----------|
| CONSTRUCTION | \$2,481K | \$6,726K | \$1,179K | \$252K | \$10,638K |
| R&D | \$484K | \$582K | \$294K | \$0K | \$1,360K |
| ENGINEERING | \$271K | \$0K | \$0K | \$0K | \$271K |
| SUBTOTAL | \$3,236K | \$7,307K | \$1,474K | \$252K | \$12,269K |

| | COST | PERCENTAGE |
|--------------|-----------|------------|
| R&D | \$1,360K | 16.76% |
| EDIA | \$2,753K | 30.59% |
| CONSTRUCTION | \$8,111K | |
| CONTINGENCY | \$4,139K | 38.11% |
| TOTAL | \$16,362K | |



**GEM Detector
Liquid Argon
EM Calorimeter
Parallel Plate**

Computed Parameters for Complete Detector

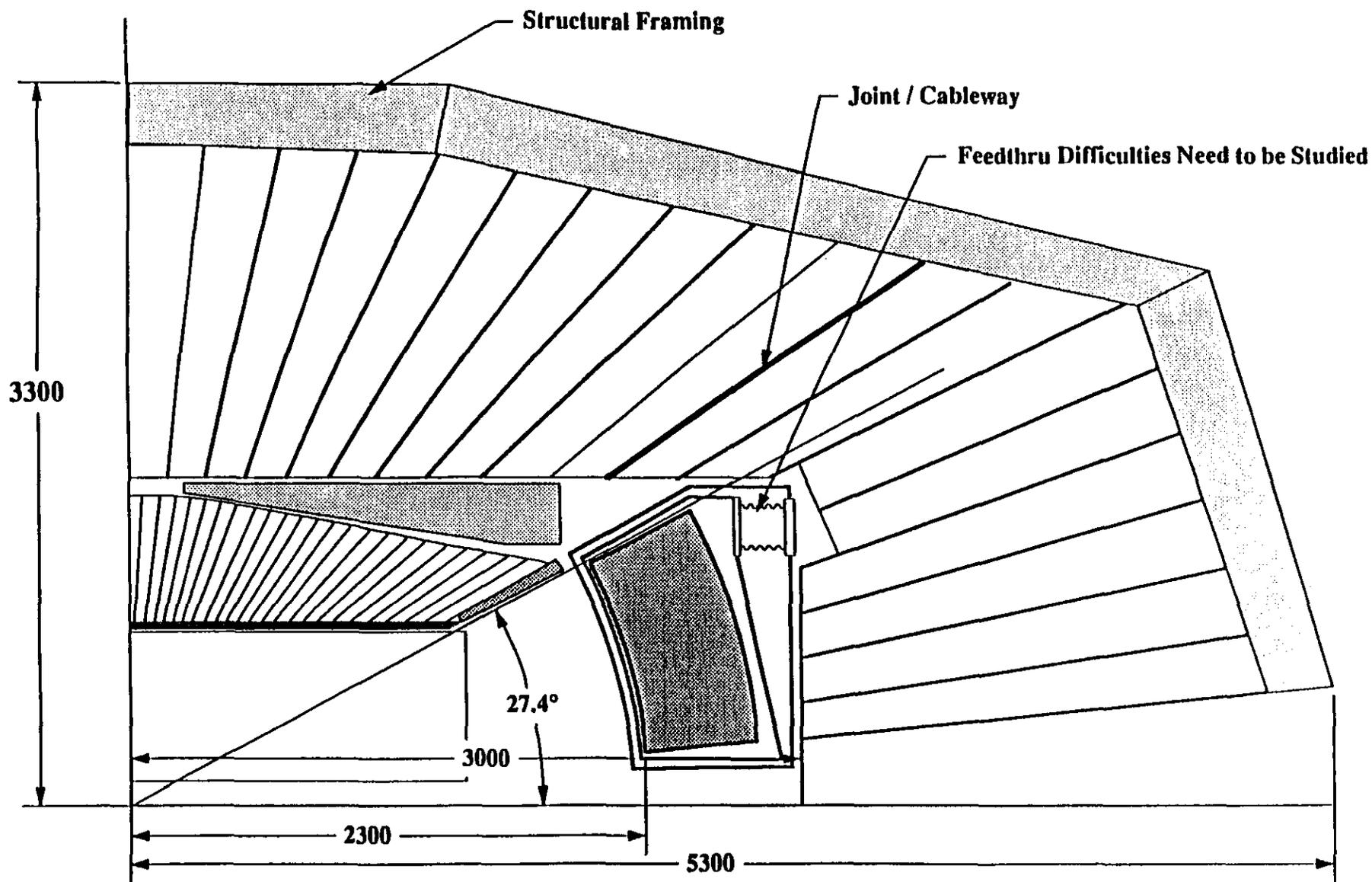
| | Totals | PreRadiator | EM Cal | Detector | |
|-------------------------------|---------------|--------------------|---------------|-----------------|--|
| Segments | 85,624 | 3,051,328 | 3,136,952 | each | |
| Channels | 31,136 | 31,136 | 62,272 | each | |
| Striplines | 1,948 | 7,784 | 9,732 | each | |
| Types of Stripline | 42 | 42 | 84 | each | |
| Length of Stripline | | | 5,357 | M | |
| Connections | 281,697 | 3,814,160 | 4,095,857 | Each | |
| PreRadiator Plate Area | 408 | 4244 | 4,652 | M ² | |
| Volume of Lead | 0.81 | 6.37 | 6.98 | M ³ | |
| Weight of Lead | 6.95 | 72.25 | 79.20 | Ml | |
| Area of Copper | 816 | 8,488 | 9,304 | M ² | |
| Weight of Copper | 0.18 | 1.92 | 2.10 | Ml | |
| Area of PrePreg | 816 | 8,488 | 9,304 | M ² | |
| Weight of PrePreg | 0.42 | 4.33 | 4.74 | Ml | |
| Area of electrodes | 408 | 4,244 | 4,652 | M ² | |
| Weight of Electrodes | 0.14 | 1.49 | 1.63 | Ml | |
| Volume of LKr | 0.73 | 7.64 | 8 | M ³ | |
| Weight of LKr | 1.25 | 12.99 | 14.23 | Ml | |
| Area of Amplifier | 31.14 | 48.51 | 79.65 | M ² | |
| Active Material Weight | 8.94 | 92.97 | 101.91 | Ml | |

Parallel Plate LKr Calorimeter
Category Distribution

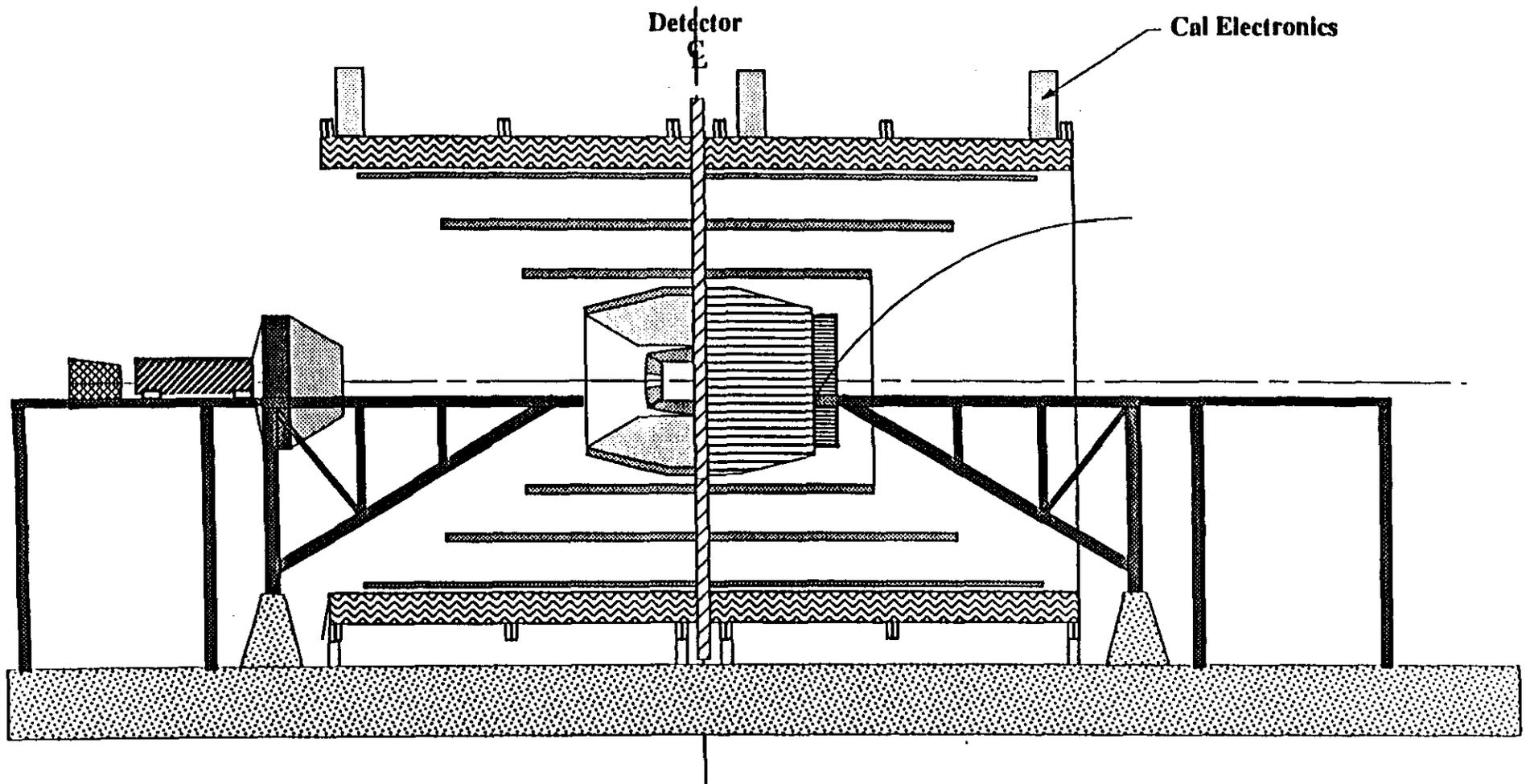
| Category | Estimate | Fraction |
|------------------------------|------------------|-----------------|
| Research & Devel. | \$2,005K | 10.3% |
| EDIA | \$6,517K | 23.9% |
| Construction | \$19,461K | |
| Contingency | \$9,766K | 38.2% |
| Total | \$37,750K | |

GEM Detector Silicon Pre-Radiator

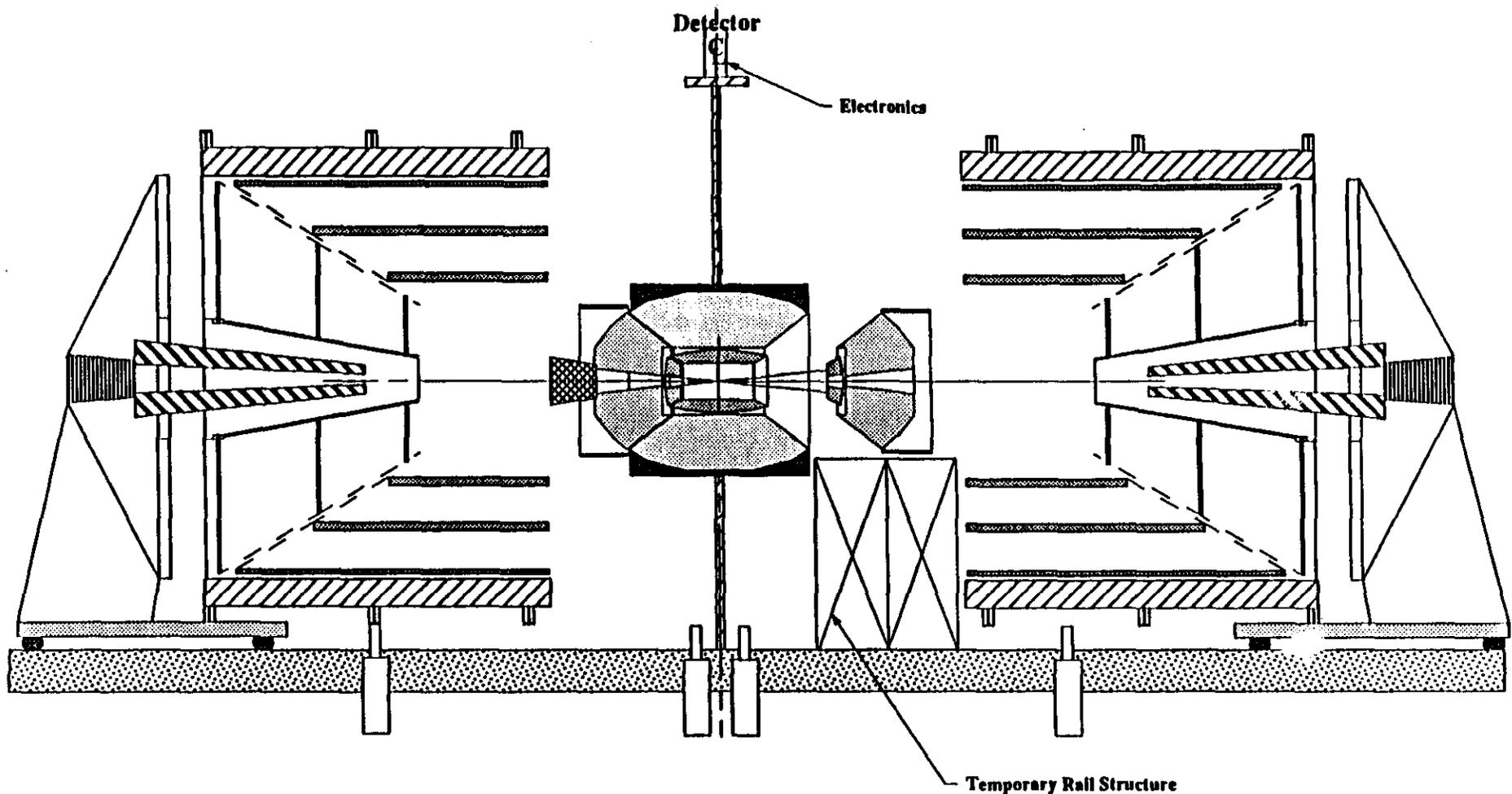
| | Catagory | Tot Uts | \$K/UNIT | Units | Cost |
|------|--------------------|---------|----------|---------|------------------|
| 1.00 | Material | | | | |
| 1.10 | Skis | | | | \$90K |
| 1.20 | Silicon | | | | \$400K |
| 1.30 | MultiLayer Cards | | | | \$338K |
| 2.00 | Readout | | | | |
| 2.10 | Electronics | | | | \$1.100K |
| 3.00 | Structure | | | | |
| 3.10 | Cylinders | 2 | 300.00 | Each | \$600K |
| 3.30 | End Plates | 4 | 100.00 | Each | \$400K |
| 4.00 | Thermal Cont | | | | \$980K |
| 5.00 | Assembly | | | | |
| 5.10 | Ass & Test | 4 | 0.045 | MH/chip | \$2.761K |
| 5.20 | Tooling | | | | \$500K |
| 6.00 | Testing | | | | |
| 6.10 | Test Beam Labor | | | | \$500K |
| 6.20 | Test Equip | | | | \$250K |
| 7.00 | Transport. | | | | \$100K |
| 8.00 | Installation | | | | |
| 8.10 | Install Labor | 5550 | 0.045 | MH | \$250K |
| 8.20 | Install Equip | | | | \$100K |
| | Direct Cost | | | | \$8,369K |
| | EDIA | 25% | | | \$2,092K |
| | Base Cost | | | | \$10,461K |
| | Contingency | 40% | | | \$4,184K |
| | Subtotal | | | | \$14,646K |
| | R&D | 11% | | | \$1,611K |
| | TOTAL | | | | \$16,257K |



**GEM Detector
BaFFILAr Calorimeter**



**GEM Configuration Option Study
Scintillating Calorimeter
Fixed Coil Version
ELEVATION VIEW**

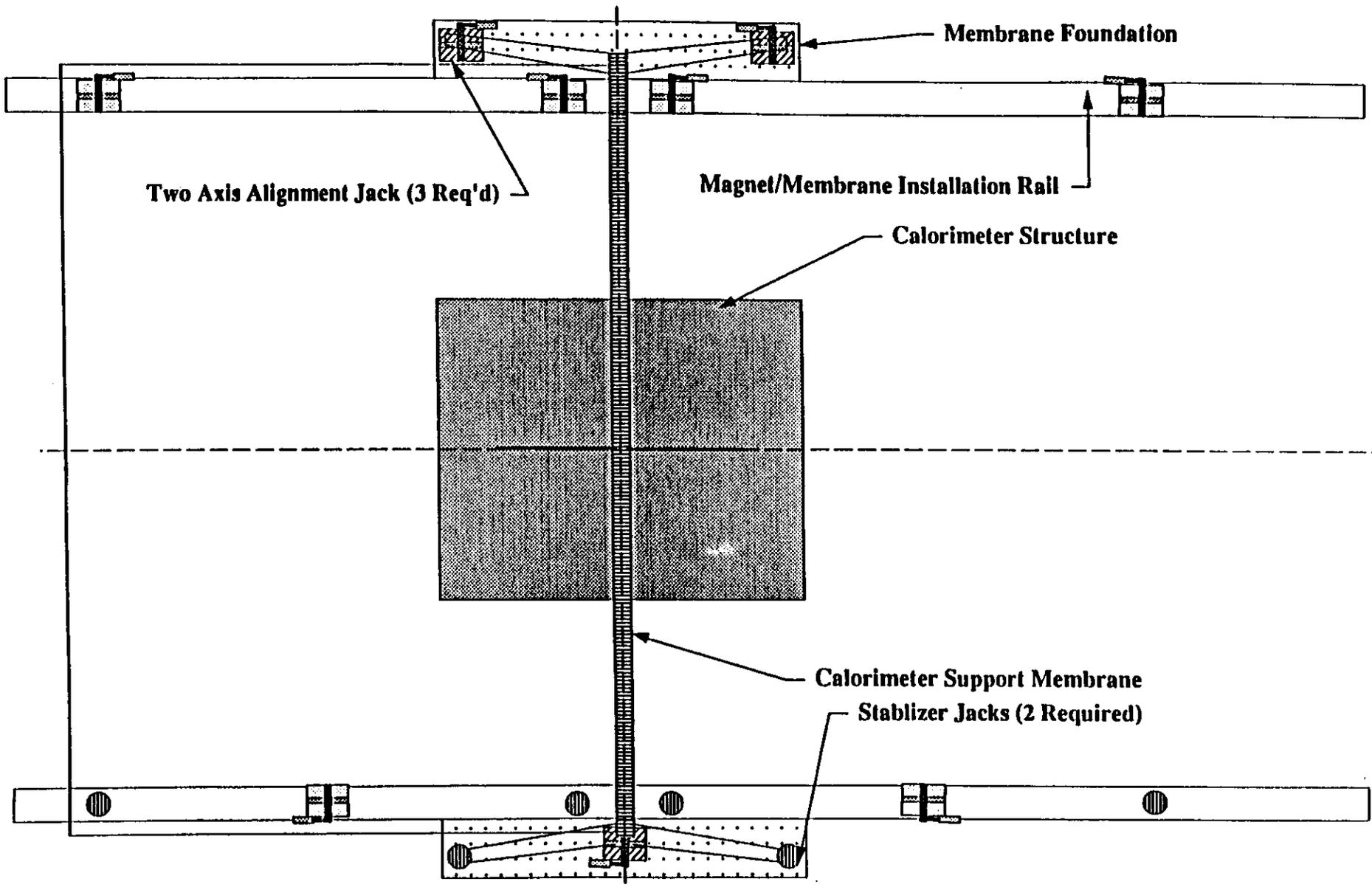


**GEM Configuration Option Study
 Scintillating Calorimeter
 Integral Calorimeter Version**

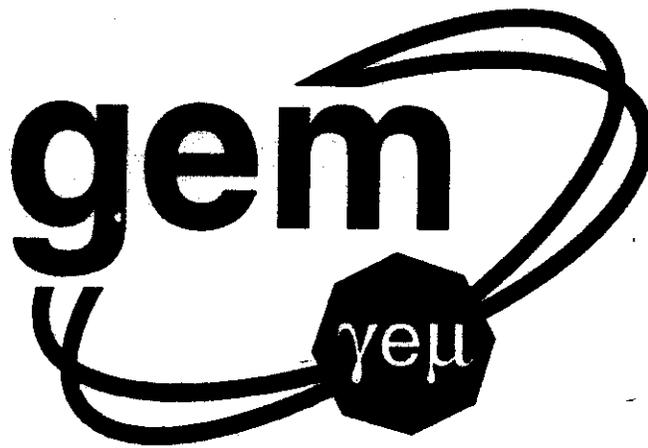
G.03.AS.00010

EBERLE/RENNICH

**Note: One Side Access Possible with Revised
 Structural Arrangement**



**GEM Detector
Membrane Support System**



Presentation by:

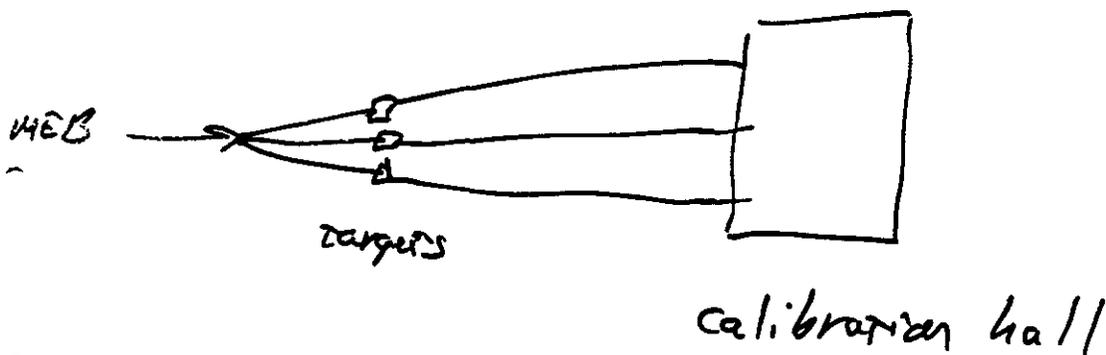
George Yost

SSCL TEST & CALIBRATION

BEAMS & HALL

3 BEAMS FROM MEB.

200 GeV primaries - we will
get secondaries.



Beam structure 16 ns bunch separation

~ 800 bunches

10^{10} p/bunch for main ring operation

5×10^{10} p/bunch for test beams operation

8 sec refill time

≤ 1 sec spill time

ping ~ few ns possible - Need requirements

max flux $\sim 10^7$ n's / pulse

anything less is easy

Serious radiation safety questions @

max intensity: 6' concrete shielding -

(Szefanski memo)

we need input on:

space req's (anything that will influence conventional construction)

safety concerns (")

utilities - power, gas, cooling, etc.

Beam flux needs

Also urgent: Needs for '92, '93, '94, ...

We must present a plan to SSCL
management in ~ 1 month.

TEST BEAM PART-II, NP8 (. 7) - NP12 (CALIBRAT)

SSC LAB

| Activity Description | Start | Finish | Dur | 01 JAN 92 | 01 APR 92 | 01 JUL 92 | 01 OCT 92 | 01 JAN 93 | 01 APR 93 | 01 JUL 93 | 01 OCT 93 | 01 JAN 94 | 01 APR 94 | 01 JUL 94 | 01 OCT 94 | 01 JAN 95 | 01 APR 95 |
|-----------------------------|-------|--------|-----|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|
| 1. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 2. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 3. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 4. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 5. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 6. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 7. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 8. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 9. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 10. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 11. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 12. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 13. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 14. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 15. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 16. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 17. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 18. Procurement of Test Bed | | | | | | | | | | | | | | | | | |
| 19. Design of Test Bed | | | | | | | | | | | | | | | | | |
| 20. Procurement of Test Bed | | | | | | | | | | | | | | | | | |

6 Apr end of July

10 Feb

10x10x10
or 10x10x10

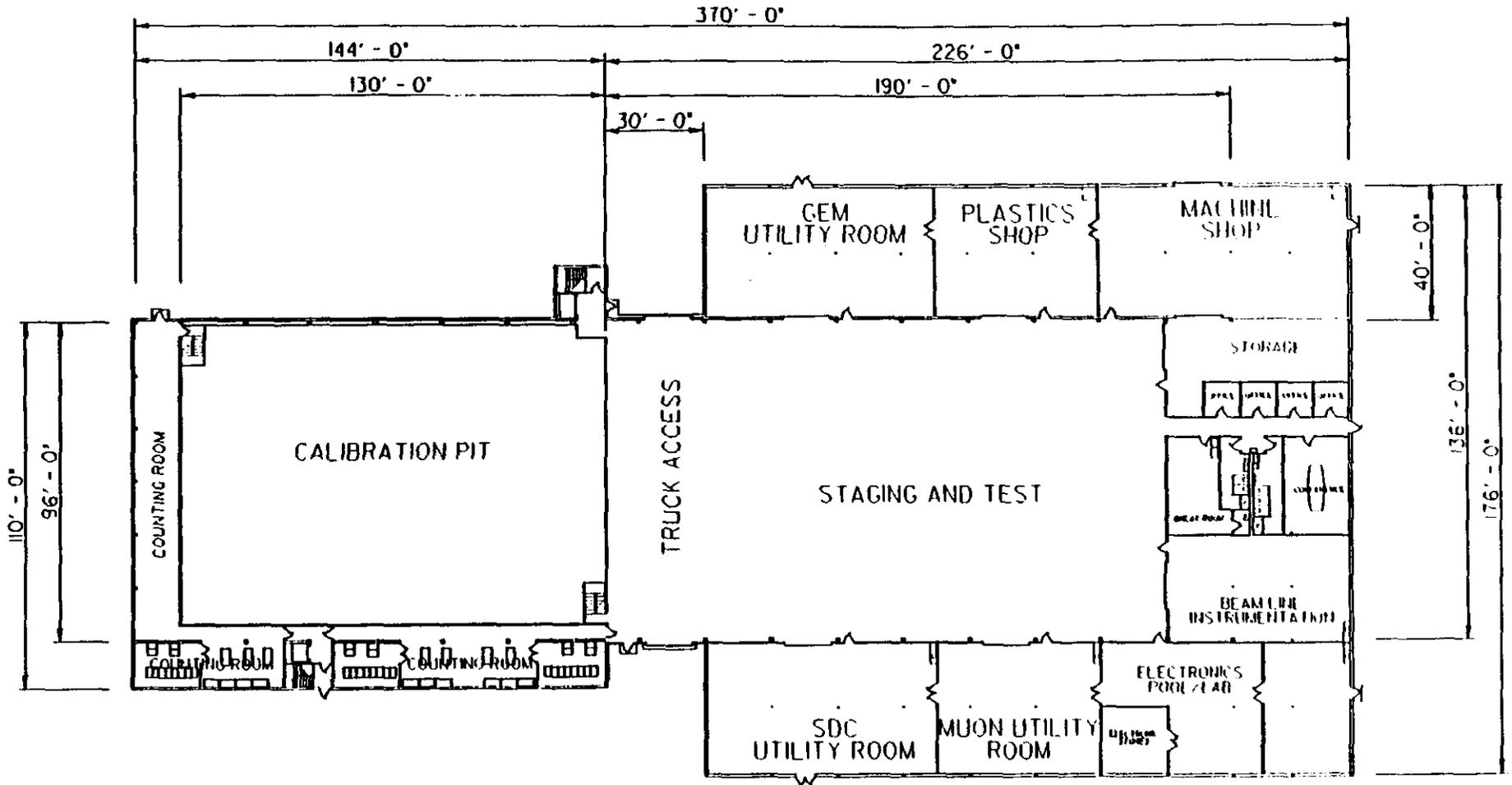
July
Dec

Bar Chart Key: Early Dates

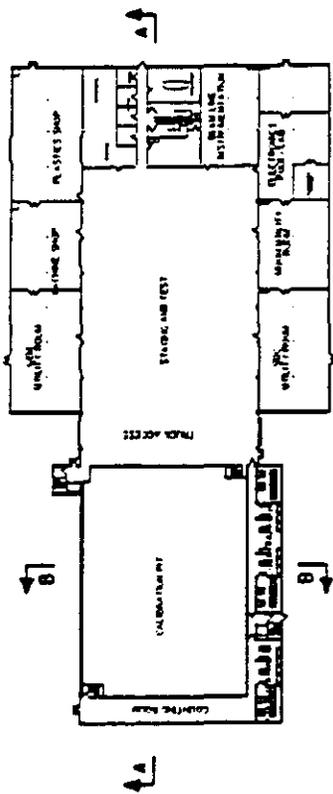
Signatures

Date:

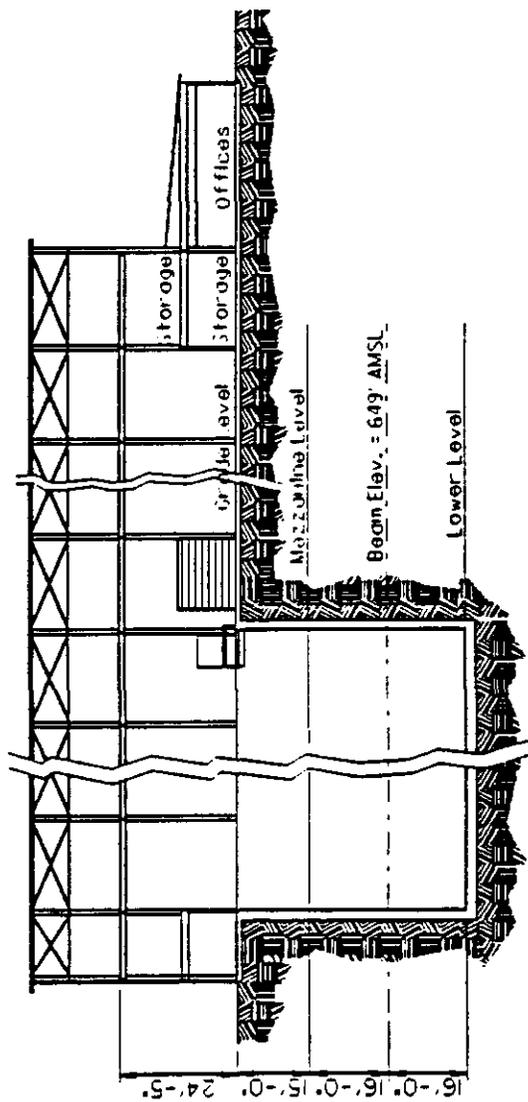
1. Name
2. Address
3. Phone No.
4. E-mail



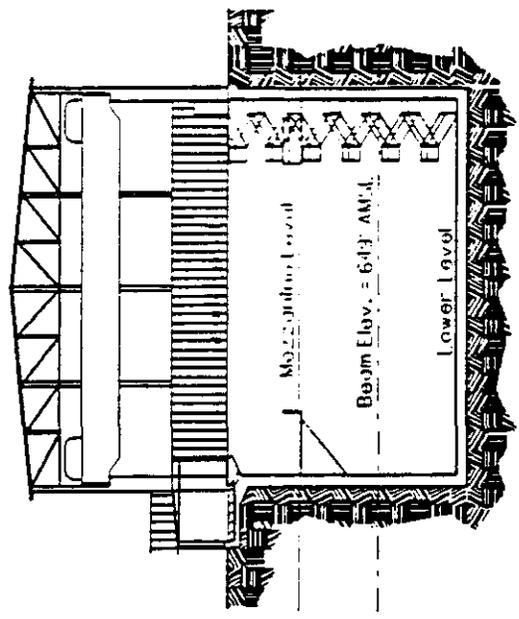
PLAN VIEW



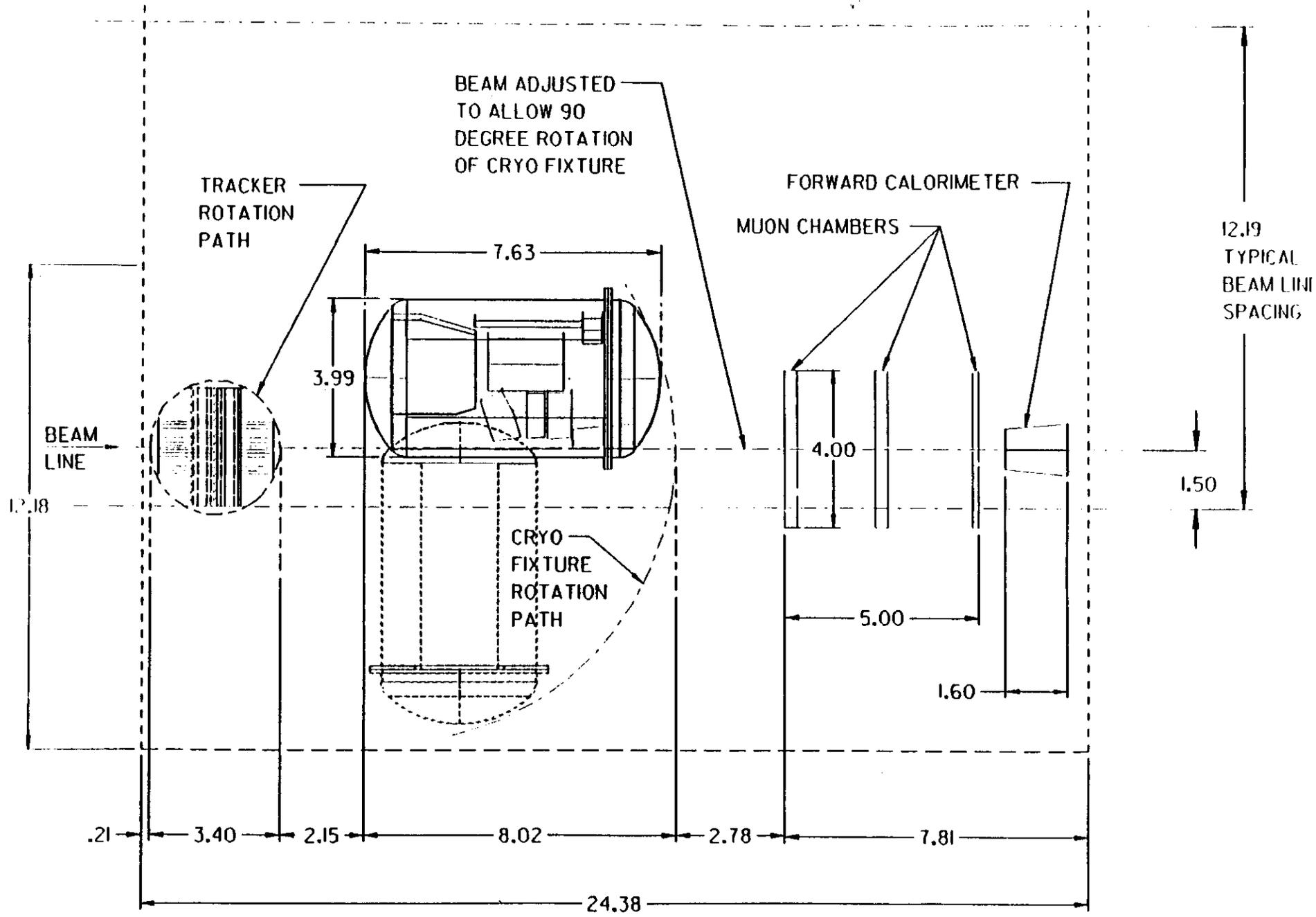
PLAN VIEW



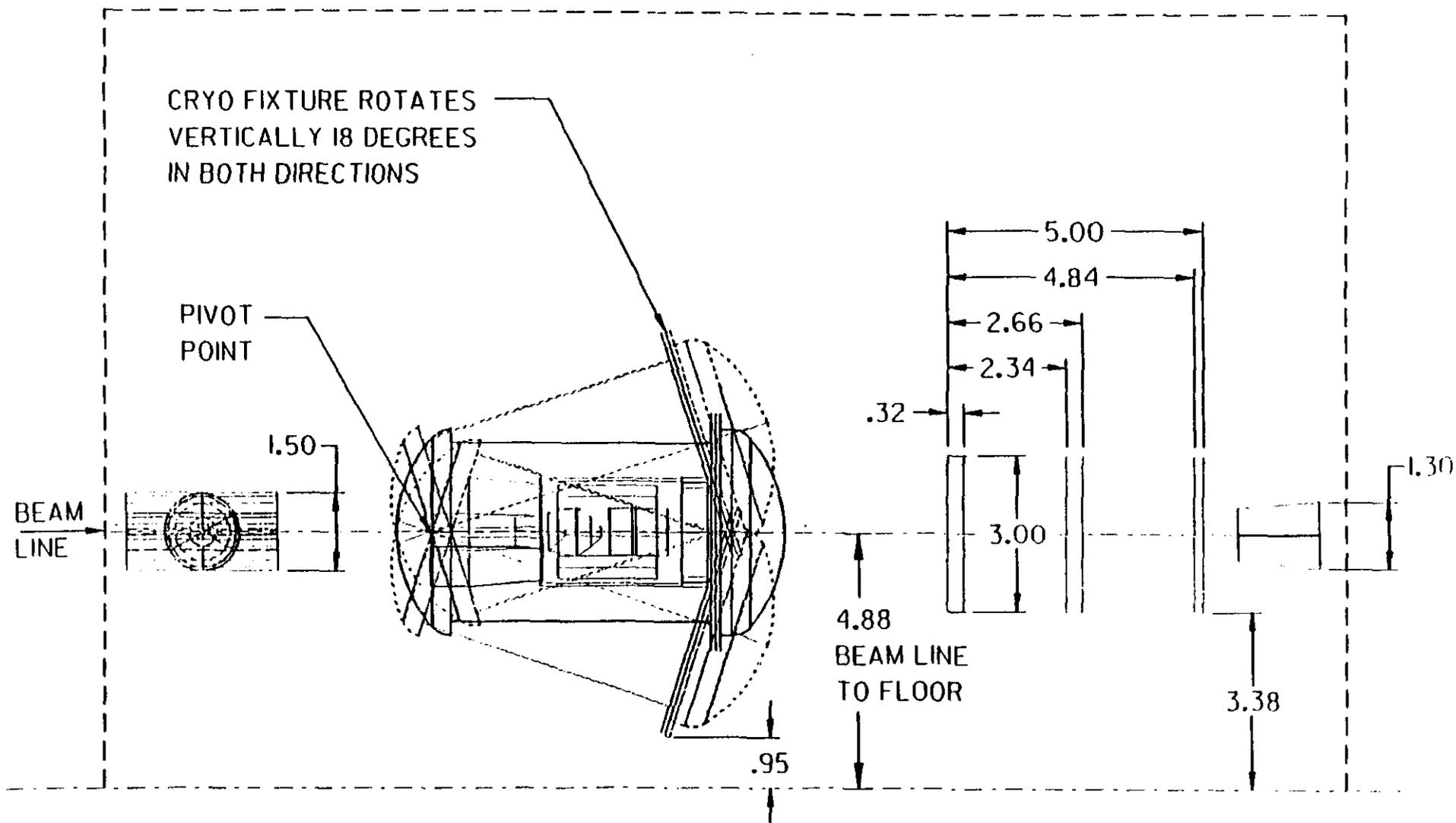
SECTION A-A



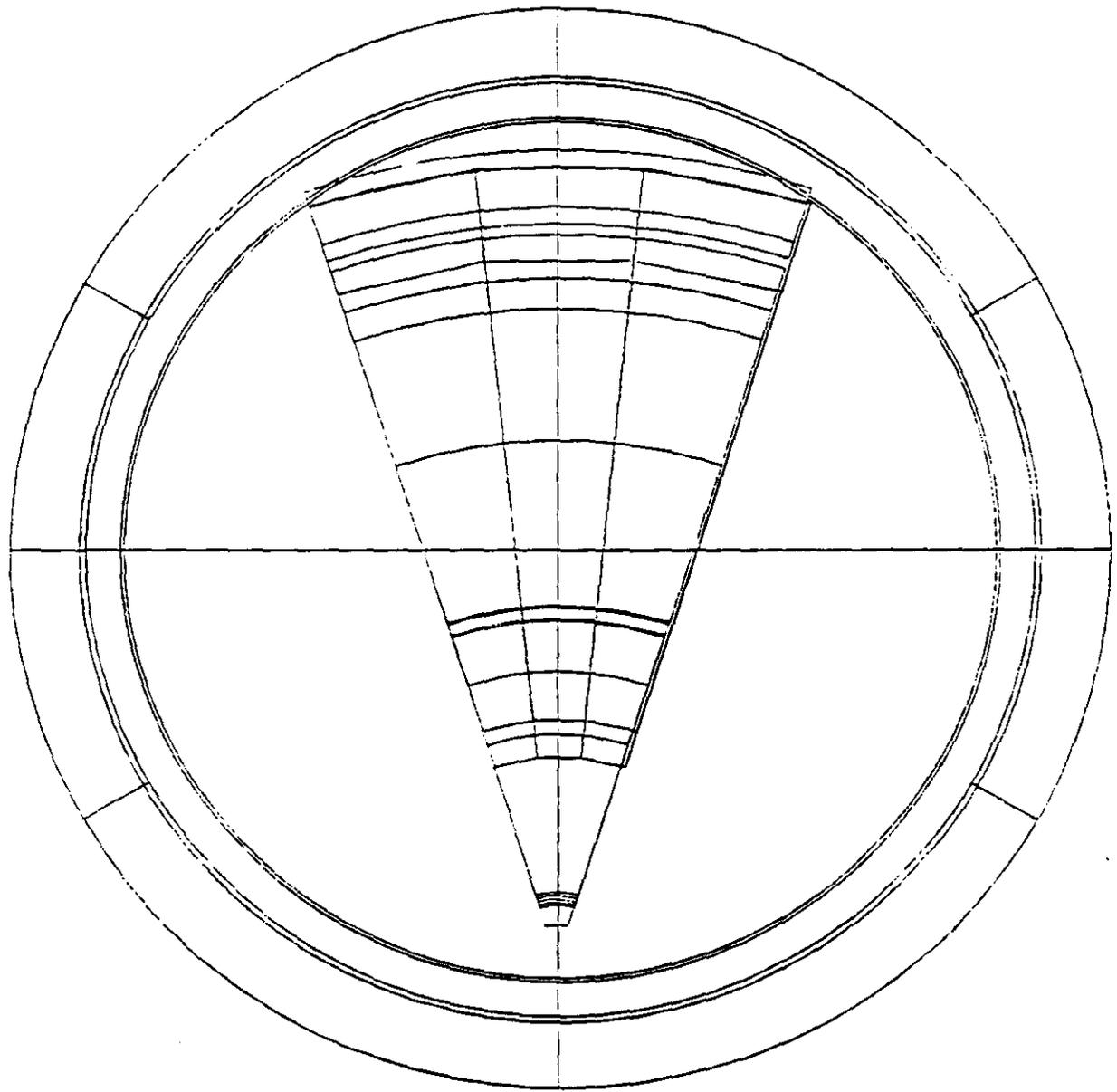
SECTION B-B

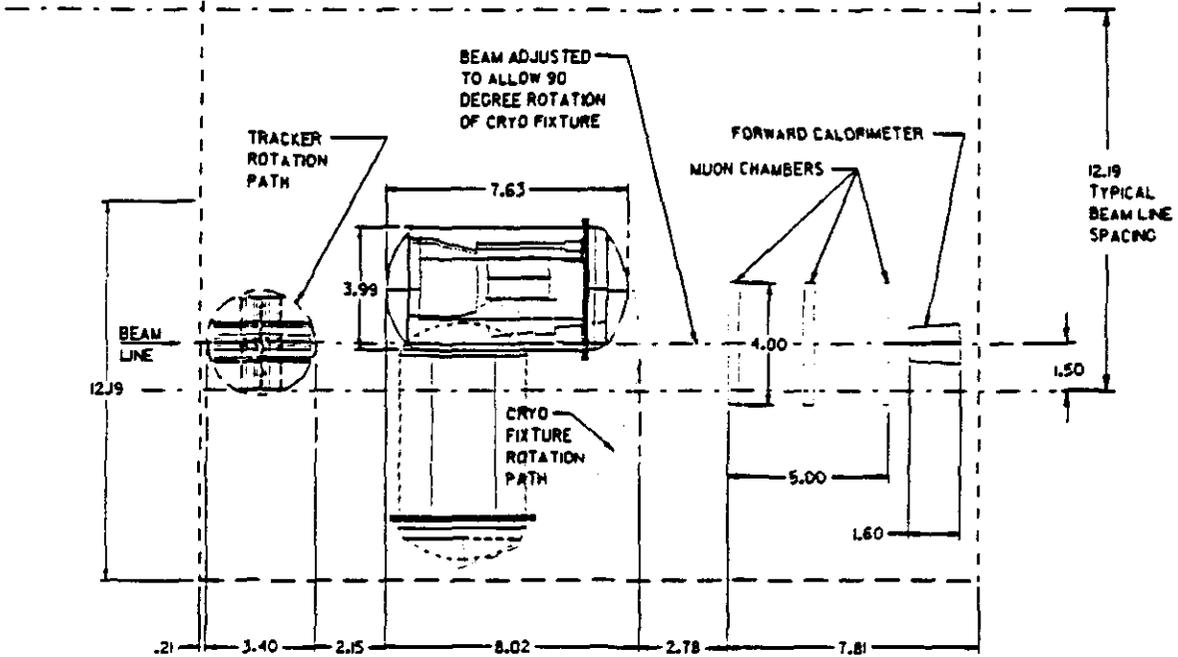
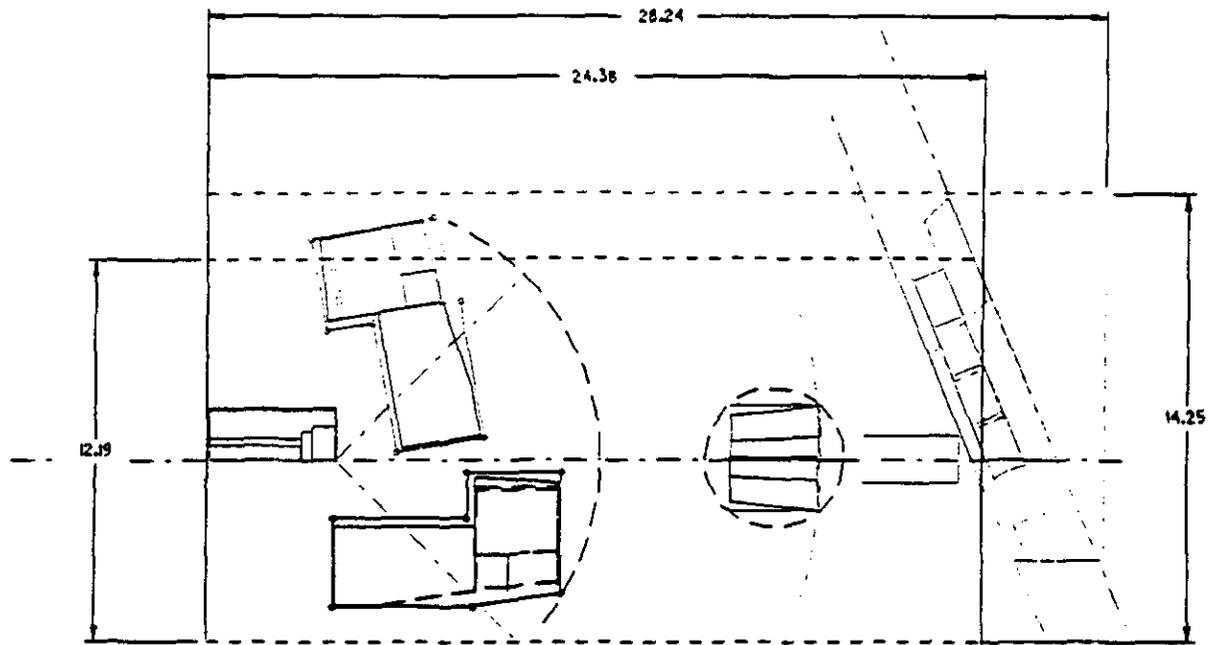


PLAN VIEW



ELEVATION





Superconducting Super Collider Laboratory
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Memorandum

To: File
From: Ray Stefanski *Ray*
Subject: Shielding Requirements for Calibration Hall
Date: Wed, Apr 29, 1992 (revised) sorry for the error!

After our meeting on test beams held in Fred's office on April 28, I consulted with Steve Butalla (at Fermilab RSO) on their general shielding requirements for test beams. We recall that the DO and CDF test-beam facilities at Fermilab have no concrete shielding associated with them. Areas that require control for radiation safety are fenced, with access through interlocked gates. This is the model that we used in establishing parameters for the Calibration Hall at SSC, namely we did not account for the possible use of shielding block in these beams.

Apparently, acceptable guidelines for radiation shielding have changed dramatically in the last two years. Steve sited an incident where a long thin detector was inserted in a test beam, without going through formal preparation, and raised radiation levels in the surrounding area by more than an order of magnitude. It's this kind of experience which has led to a policy at Fermilab to provide shielding to account for the worst event that could possibly occur in the area. Administrative or barrier controls are no longer acceptable.

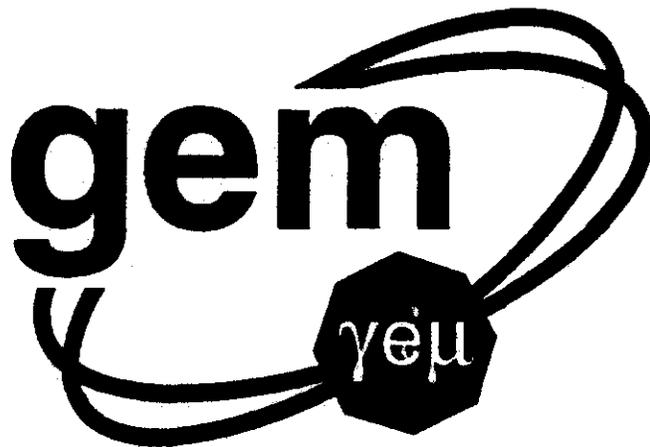
Steve did a back of the envelope estimate of shielding requirements for operation in a hadron beam at 10^7 particles per second and a flat-top of one second in an eight second cycle time. He sited FNAL-TM 1140 as a reference for shielding guidelines. (I've asked the Fermilab library to send me a copy.) The assumption is that one must provide shielding for a thin target, three interaction lengths in length (one assumes that we achieve shower maximum in the target), and that the occupied area is about six meters from beam center. The occupied area must be kept below 0.1 mrem/hr, which is conservatively set below the guideline of 0.25 mrem/hr established by DOE for such areas. The result is:

Two meters of concrete for 10^7 particles per second,
One meter of concrete for 10^6 particles per second,
and 2.5×10^4 particles per second for no shielding.

One can assume that radiation levels fall as the inverse of the square of the distance for distances beyond six meters.

These requirements would apply to areas where people are expected to work for extended periods. This would not apply to catwalks, for example, since occupancy would be for very short duration. Shielding for the roof of the building would also probably not be required.

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Ronn Woolley Bob Lavelle Tom Murphy
Howard Fenker Frank Stocker Steve Butalla(FNAL)



Presentation by:

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Barium Fluoride Surface Preparation and UV Coating R&D at LLNL - A Progress Report

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1. Introduction

Lawrence Livermore National Laboratory (LLNL) has been asked to provide R&D on new methods of barium fluoride (BaF_2) crystal surface preparation, as well as optical coating methods to improve the efficiency of transport of scintillation light through the crystal to the photodetector. The goal of the R&D is to identify a method (or methods) for preparing the surface to eliminate the amorphous layer that has been found to be a product of the surface preparation techniques used when the crystals are manufactured in China and provide a crystalline surface free from oxygen and other contamination, with suitable flatness for optical coating. UV reflective coatings have been identified that are suitable for the wavelengths of the scintillation light (220 nm peak). The coating consists of a layer of aluminum with a magnesium fluoride over-layer. This coating can be applied in a uniform layer or in a graded layer, with reflectance varying as a function of position. It is ultimately desired that the UV coating be applied with graded reflectance to linearize the response of the crystal along its length.

2. BaF_2 Surface preparation R&D at LLNL

A number of surface preparation techniques have been identified as potentially capable of removing the amorphous layer and providing a clean crystalline layer for coating. The R&D program, which began in early March, 1992 has explored the following surface preparation techniques:

A. Polishing with diamond grits

BaF_2 crystals have been polished at LLNL using this standard technique for use as laser windows. Up to this point it has not been known how the crystal structure is affected. Comparison of this polishing technique with diamond turning has shown that diamond turned laser windows are less susceptible to damage in high intensity laser beams. The process uses diamond impregnated pads and a BaF_2 sample was polished to 20 angstrom RMS surface finish for study. This technique is probably not suitable for large area polishing because edges tend to be rounded off and flatness over large areas is not easily achieved.

B. Ductile grinding

The process known as ductile grinding uses a specially prepared wheel coated with diamond. The wheel is prepared so that it is flat to within 0.25 - 0.5 microns and the part to be polished is translated across the surface of the rotating wheel. In effect, this is using many diamond turning bits instead of one. The advantage of this process is that it is very suitable for mass production polishing of large numbers of crystal surfaces. LLNL's ductile grinding system was dismantled last year and I have requested that it be rebuilt. We do not have results from this process yet.

C. Diamond pitch lapping

This process uses specially prepared wheels coated with a synthetic pitch and impregnated with diamond powder. The prepared wheel is rotated and the sample to be polished is translated back and forth across the wheel as the wheel rotates. The sample is usually immersed or heavily washed with a low viscosity silicon oil. We have prepared a small sample using this technique and have measured a surface finish of about 10 angstroms RMS. This technique has the advantage of being simple to implement and can provide large area and large quantity polishing capability.

D. Diamond turning of the crystal surface

Diamond turning of crystal surfaces are routinely done at LLNL in support of the Laser Program and other programs requiring precision optical surfaces. LLNL's Large Optics Diamond Turning Facility (LODTM) is capable of turning surfaces with radii as large as 1.65 meters with a precision at the few angstrom level. Other smaller diamond turning machines are available for preparing surfaces with similar precisions.

We opted to perform "wet" diamond turning using a highly purified water-free lubricating fluid (Drakeol 7, a low viscosity poly-alpha olefin oil). Alternative cutting fluids include low viscosity silicone oil or a pure hydrocarbon such as hexanes. Drakeol 7 is routinely used for diamond turning potassium di-hydrogen phosphate crystals (KDP) for use as frequency doubling and tripling crystals for the Nova laser at LLNL. The crystals are turned and then cleaned using a pure toluene soak for 24 hours. A similar cleaning method was used on BaF₂ crystals.

We achieved very successful results using small samples of BaF₂ crystals. In one sample we were able to achieve a 6 angstrom RMS surface finish measured over a 600 micron area of the crystal. We also observed that the surface finish depended on the crystal lattice orientation. In one sample we observed markedly different surface finish quality when machining across a crystal grain boundary. The surface finish on one side of the boundary was found to be 80 angstroms RMS compared with 6 angstroms RMS immediately on the other side of the grain boundary.

In summary BaF₂ appears to be a good material for diamond turning and other diamond cutting processes, such as ductile grinding and diamond pellet lapping.

E. Argon Ion Beam Milling

Argon ion beams are also available for removing material from the crystal surface. In this technique a beam of argon ions is directed onto the surface to be milled, with the rate of material removal determined by the orientation of the crystal surface with respect to the beam. For our studies the crystal was oriented at 45 degrees. Removal rates vary depending on the material but typically are a few microns per minute to a few microns per hour. For our study a carbon mask about 2 cm x 1 cm was prepared to allow the crystal to be milled in a controlled manner. Carbon is a good material for the mask because of its extremely low sputtering rate compared to most materials.

We prepared three samples with measured removal depths of about 0.5, 1.0 and 2.0 microns. The samples took about 1/2 hour to remove 1 micron of material. Surface quality was observed to be poor (compared to previously described surface finishes) and in general followed the original surface quality.

2. X-ray diffraction analysis and energy dispersive spectroscopy

Crystals were analyzed, before and after preparation using x-ray diffraction analysis, and electron beam microprobe methods for the determination of structural and chemical properties of the surface, respectively. We noticed that x-ray diffraction measurements would show the BaF₂ lines slightly shifted and additional unidentified lines would also be present. Unfortunately, we were unable to precisely identify the nature of these spurious lines. Also, x-ray diffraction was unable to qualitatively distinguish between crystalline and amorphous surfaces. We did closely examine the sample crystal that exhibited the two different surface finishes on diamond turning. We cleaved small samples from the two regions and crushed them into a powder. There was no noticeable difference in the two materials.

We also performed energy dispersive spectroscopy (EDS) on a sample and found the strongest signals due to strontium and iodine.

3. Rutherford Backscattering Analysis at Charles Evans & Associates

Three BaF₂ samples were brought to Charles Evans and Associates for Rutherford Backscattering (RBS) analysis. The samples were: 1) the diamond turned piece with the two different surface finishes, 2) the diamond pad polished piece with the 20 angstrom surface finish, and 3) the ion beam milled sample with the three different depths of cut.

Measurements were first made on sample 1) for the two different finish regions. No channeling was observed on either area for this sample, leading us to believe that the crystal surface on the sample is amorphous

The second sample measured was the ion milled sample 3) which had to be broken down to a smaller size to fit in the RBS machine. We opted to look at the 2 micron depth milled portion. In this measurement, strong channeling was observed, indicating a crystalline layer had been exposed. In addition RBS was performed across the sample on a portion of original Chinese surface finish that was masked off. In this region no channeling was observed.

The third sample measured was the standard polished sample 2). Measurements are still in progress on this sample.

A fourth sample, prepared using the diamond pitch polishing method is being measured at this time.

4. UV Reflective Coatings

LLNL regularly coats optics with both reflective and anti-reflective coatings. Magnesium fluoride is typically used for anti-reflective coatings. The reflective coating to be applied to the BaF₂ crystals consists of a thin layer of aluminum (about 500 angstroms) with a MgF₂ over-layer. Reflective coatings of this type exhibit high reflectance, typically 75%-85% down to about 180 nm. In addition the coatings are robust. We utilize an electron beam evaporative coating system specifically designed for the purpose of laying down MgF₂ and aluminum.

We have prepared two different samples for the purpose of characterizing the quality of the reflecting coatings. One sample was a large piece of BaF₂, left with the original Chinese surface finish, about 6 inches long cut from a larger prepared crystal pair. The other sample was a small BaF₂ sample with a high quality surface finish for use as a witness sample. This witness sample was used to quantitatively characterize the quality of the reflective coating.

Measurements were made on the witness sample for both front surface reflectivity and back surface (through the BaF₂) reflectivity. The front surface reflectivity was measured to establish that the coating was of the proper quality. Results of the front surface measurement confirmed that the coating was high quality, with a reflectivity of about 90% from 300 nm to about 220 nm followed by a gradual drop to about 86% at 190 nm. Back surface reflectivity showed that the reflectance was about 87% at 300 nm falling gradually to about 85% at 240 nm with a steeper fall to 74% at 190 nm. At 220 nm the reflectivity is about 81%.

A qualitative look at the large BaF₂ sample shows that surface finish plays an important role in the uniformity of the coating. Pin hole sized areas were observed that were not coated, presumably due to the observed scoring of the crystal surface during the Chinese surface preparation.

These initial samples were coated with uniform reflectance coatings. Gradient coatings will also be studied. Two alternative ways of performing gradient coatings exist. One involves translating the sample past the source at a varying rate. The other involves placing the sample at an angle with respect to an isotropic coating source. It is likely that the translation method will be adopted because it is suitable for long crystals and translation stages are easily programmed for variable motion to allow the deposition of coatings with complex reflection gradients.

The Monte Carlo program LTRANS³, developed at the Institute for Theoretical and Experimental Physics (ITEP), Moscow is being adapted to model the BaF₂ crystal properties. This Monte Carlo code combines scintillation light emission, crystal transmission, absorption, wall effects and geometric ray tracing to allow an evaluation of the uniformity of light transport to the end of the crystal as a function of position on scintillation light emission. This code can be used to estimate the correct gradient for the surface reflective coating to provide the optimum uniformity along the crystal length.

4. Conclusions

We have performed R&D on four different BaF₂ surface preparation techniques. All the techniques have demonstrated surface finishes better than 20 angstroms RMS. Three techniques: diamond turning, ductile grinding, and diamond pellet lapping hold promise for mass production of large area high quality surface finishes on BaF₂ crystals. Unfortunately, RBS studies show that diamond turning and standard diamond polishing does not provide a crystalline surface.

It is unknown at this time if the surface finish starts out crystalline and then becomes amorphous over time. It has been suggested that Charles Evans & Associates perform an RBS on a freshly cleaved BaF₂ sample and then follow up a few days later to see if a change in the surface is evident.

Ion beam milling is effective for removing BaF₂ surfaces down to a depth of a few microns/hour. In addition, there is an indication that ion milling provides a somewhat crystalline surface. Adapting this technique to mass production is feasible, but the quality of the surface is only as good as the surface one starts with and, in general, is worse than surfaces prepared using various grinding or machining techniques.

Optical coatings for high reflectance in the UV (> 80% at 220 nm) appear to be achievable using standard E-beam evaporative coater. We also intend to study the effect of applying a hard silicon dioxide overcoat to protect the coatings from damage.

Finally, it is worth reiterating the previously mentioned work comparing laser damage to optical windows prepared using standard polishing techniques and diamond turning. It is likely that this is a surface or near-surface phenomena and is related to stress build-up in the crystal during the polishing technique. Acid etching of surfaces reveal these stresses as micro-cracks, or pits, and acid etching is commonly used as an analytical tool in characterizing laser glass surfaces.

We are currently studying the character of BaF₂ surfaces using acid etching. For example, a sample of BaF₂ was pitch polished to 10 angstroms RMS surface finish and then one half was dipped into an acid solution consisting of 10% hydrofluoric acid, 5% hydrochloric acid and the remainder, water. After 5 minutes the sample was removed and the surface was studied under an optical microscope and a profilometer. The optical observations show small linear scratches indicative of shear stress on the crystal, however the size of the scratches indicates that the stress is small. Also, no pits are observable. The etched surface shows a 35 angstrom RMS finish and appears identical to the adjacent un-etched surface to the naked eye. We have sent this sample to Charles Evans and Associates for RBS analysis of the two surface treatments.

Despite the lack of channeling observed in RBS measurements of some of the samples we have prepared we feel that it is probably worthwhile to explore the radiation hardness of BaF₂ crystals prepared using the various techniques described in this paper.