## MIMOSA 23 SETTINGS

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## Current status

- Detector in use: FoCal prototype; 24 layers , 96 chips (MIMOSA 23)
- Non uniform response: from the testbeam data analysis we noticed that we have large variation in sensitivity from chip to chip
- Settings in use: settings used during the test beam: Vref2=80 Vref1 optimised per single chip
- Tests in lab:
- "pedestal" measurements -> systematic studies in Vref1 and Vref2
- Cosmic data taking (for given threshold settings to study the MIP response)
- Only digital R/0 available (no voltage reading)


## Open points (1)

1. Impossibility to get the discriminator transfer function normalised from 0 to 1

Fraction firing pixels (new definition) chips 8-11, Vref1=SPS-settings


## Open point (2)

- The normalisation of the firing pixels to 1 hides possible pathological behaviour


## Fraction firing pixels chips 8-11, Vref1=SPS-settings



## Open point (3)

Different Vref1 settings -> different discriminator transfer functions

## Fraction firing pixels chip 8



## Open point (3) another example

Fraction firing pixels chip 11


## Open point (4): unbalance left-right



All the chips have Vref2=80: unexpected unbalance!!!
Some unbalanced chips present hot columns/rows/pixels

## Open point (4)

- Scan in Vref2 looking at the unbalance left-right


## Balance of chip 8



## Open point (4) another example

- Scan in Vref2 looking at the unbalance left-right

Balance of chip 9


Which is the procedure to get the best Vref2 value then?

## Open point (5)

- The discriminator transfer function fit


Question: can we do the same operation but with the digital readout?

## Open point (6): clustersize (chip 8\&9)



Corrected response profile for chip 9


## Open point (6): clustersize (chip 10\&11)




## More questions might come....



