



# **Glassy metal mirror suspensions:**

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## **an update**

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**19th of March 2003**

**LIGO-G030085-00-D**

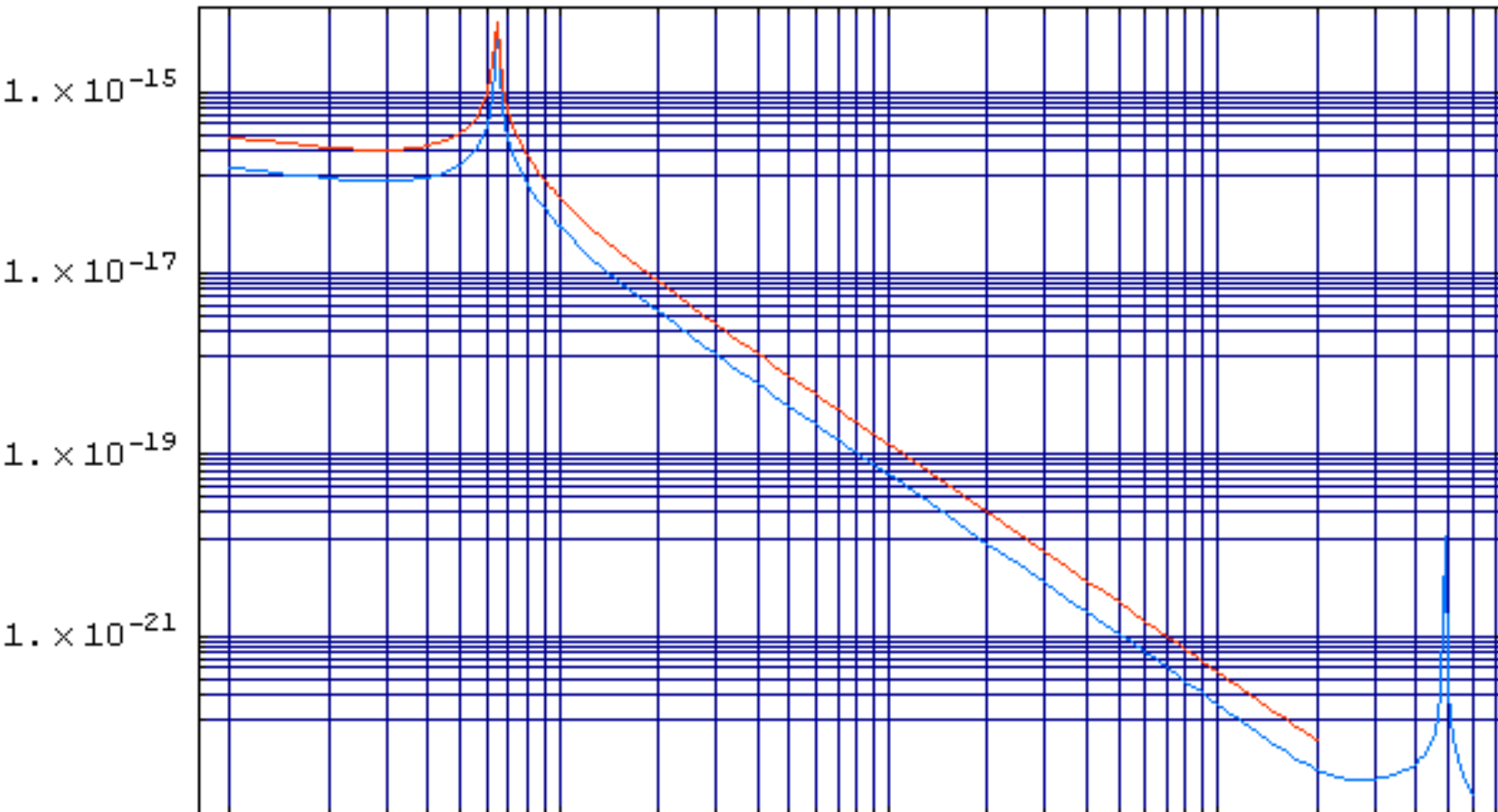
# Estimated MoRuB glass properties

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- Mo<sub>49</sub>Ru<sub>33</sub>B<sub>18</sub> in atomic percent.
- density, 9.5 g/cc
- heat conductivity, 10 Watts/m-K
- heat capacitance, 30 J/mole-K
- linear thermal expansion coeff.,  $5-6 \times 10^{-6} \text{ (K}^{-1}\text{)}$
- elastic modulus, 250 GPa
- Poisson modulus, 0.36-0.38
- breaking point  $\sim 5 \text{ GPa}$

- - These numbers should be accurate to +/- ~20%

# Thermal noise of MoRuB flex joints



Glassy metal  $Q=10^4$ , Fused  $\text{SiO}_2$  dumb bell shaped fiber  $Q=8.4 \cdot 10^8$ ,  
 $10 \cdot 3000 = 30,000 \mu\text{m}^2$ ,  $357 \mu\text{m}$  diameter,  $100,000 \mu\text{m}^2$ ,  
 60 Kg mirror, 40 Kg mirror

# Measurements of $Q$ measurement on vitrelloy

Vitrelloy has  $T_g$  of  $650^\circ\text{C}$   
while  
MoruB  $T_g$   $1050^\circ\text{C}$

MoRuB does not  
accept hydrogen

MoRuB should have  
Less important  
loss mechanisms

# Measurements on MoRuB

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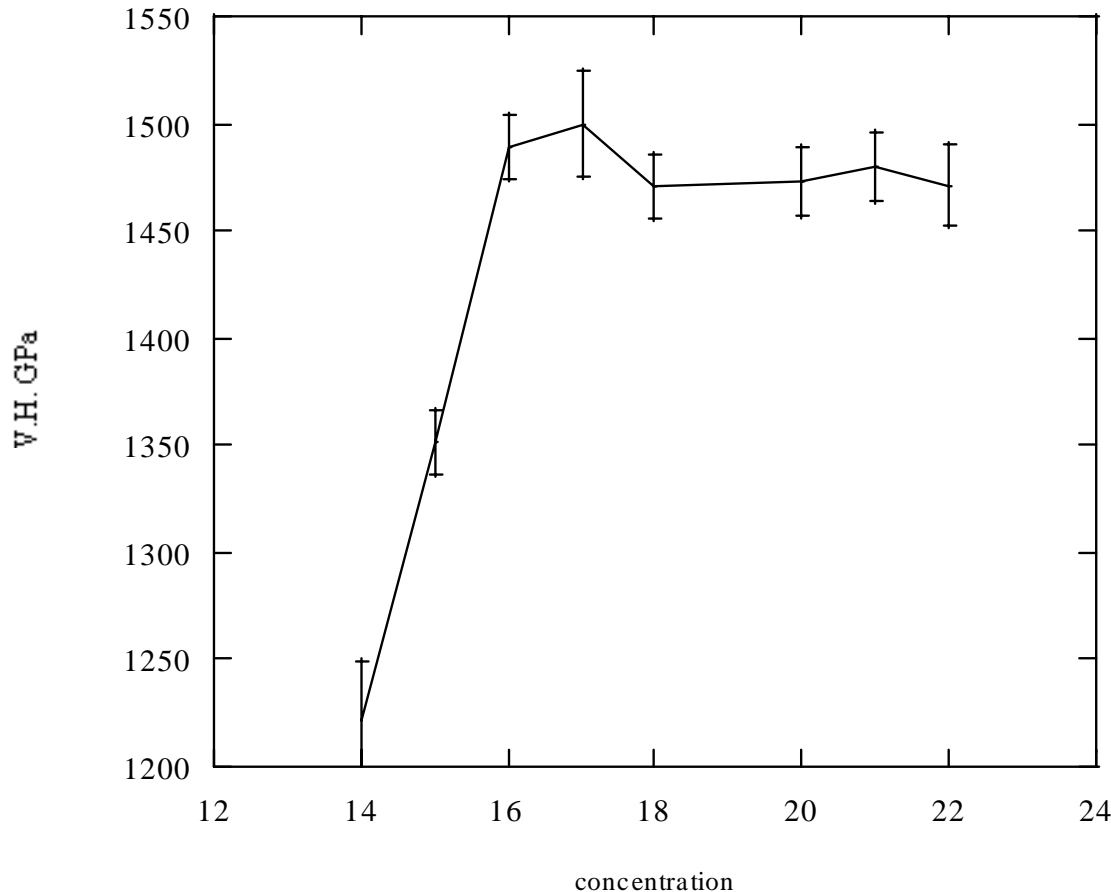
- On thick vitrelloy used modal suspension technique
- MoRuB cannot yet be manufactured in thick slabs
- Need diving board measurements
- So far with MoRuB found erratic results
- Different for each mode
- Indicating problems on the clamping point

# Identified problem with the AuSn braze

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- Found that AuSn braze, unlike standard brazes, it wets without sticking, elastic oscillations or strain detach the edge point
- S-bond, a commercial braze based on (AgSn)TiCd wets and sticks.
- But softer
- Converting to S-bond
- Discussion later

# Vicker Hardness behaviour old measurements



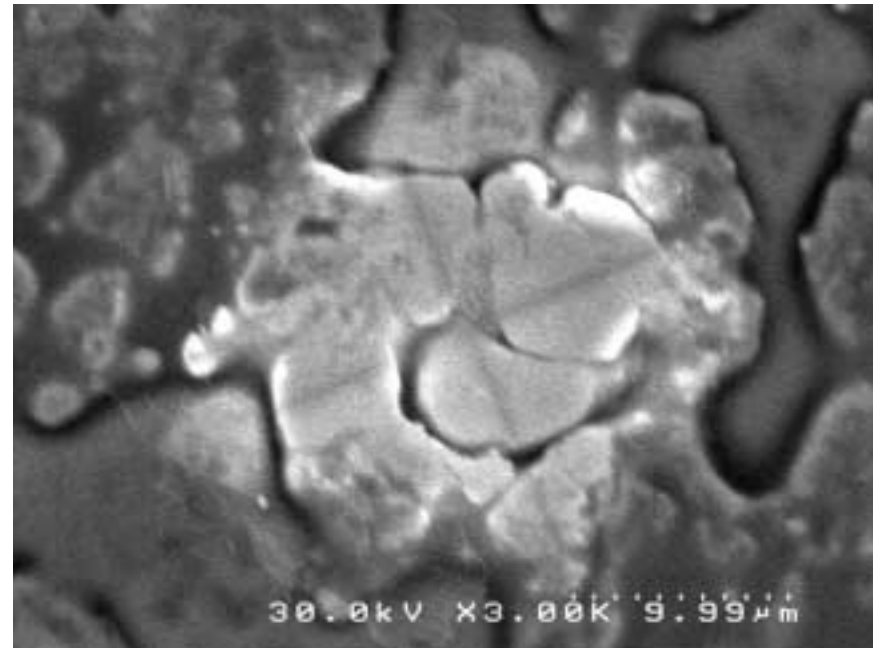
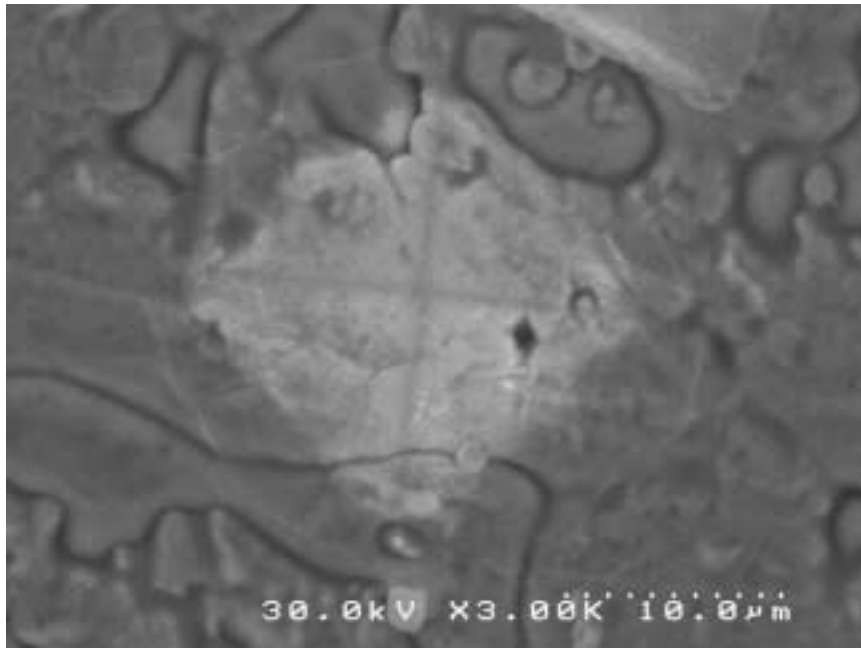
- from 14% to 16%B  
**linear behaviour**
- from 16% to 22%B  
**almost constant**

# Vicker Hardness behaviour old/new measurements

Measurements of hardness based on optical measurements of the size of indentations (see below)

Quite subjective (different people's eyes have different judgments) leading to large systematics

Large fluctuations of errors => large error bars





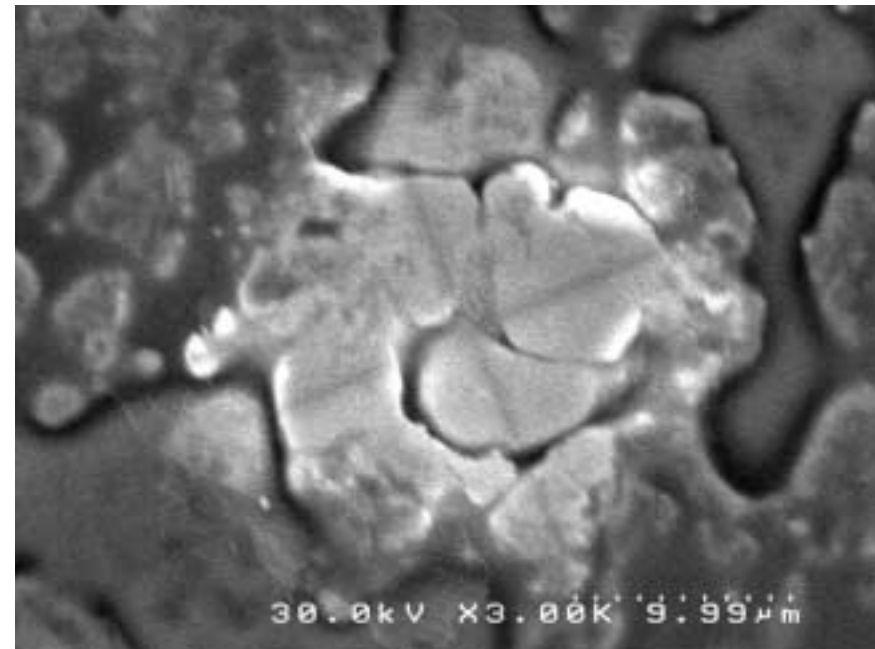
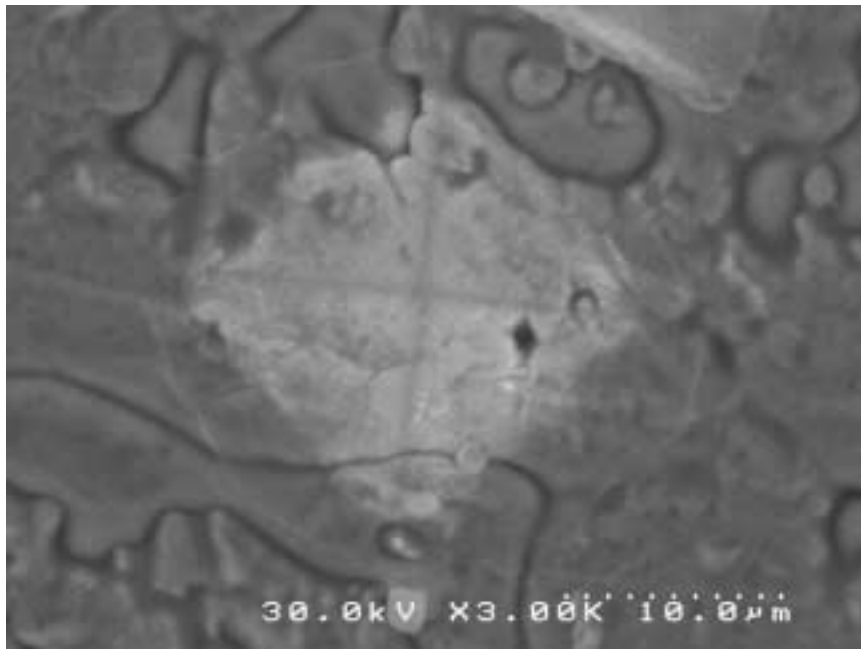
# Vicker Hardness behaviour old/ new measurements

In SEM it is obvious that the optical measurement did not even “see” the surface imperfections

⇒ Polished samples before indentation

⇒ Use SEM for evaluating indentation size

Allyson Feeney



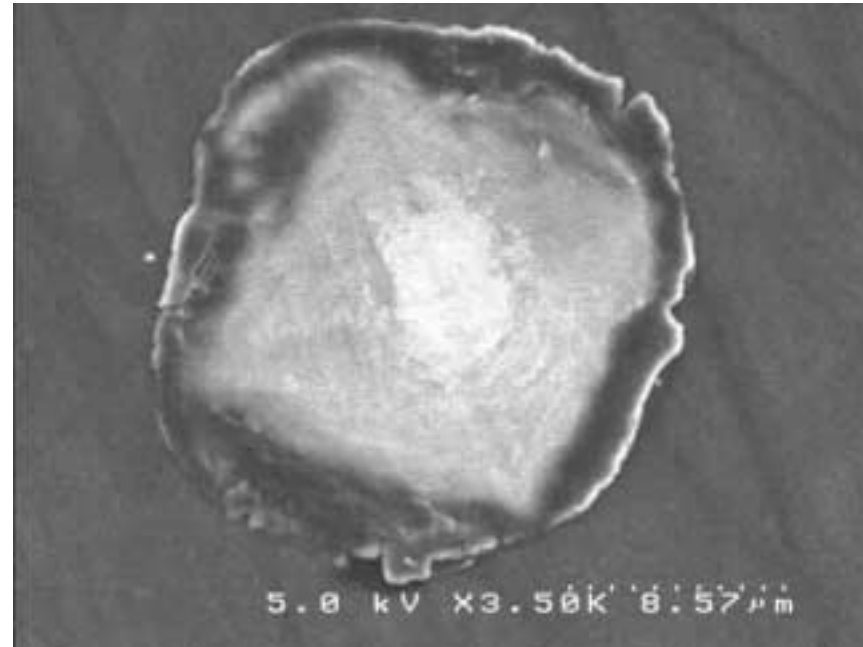
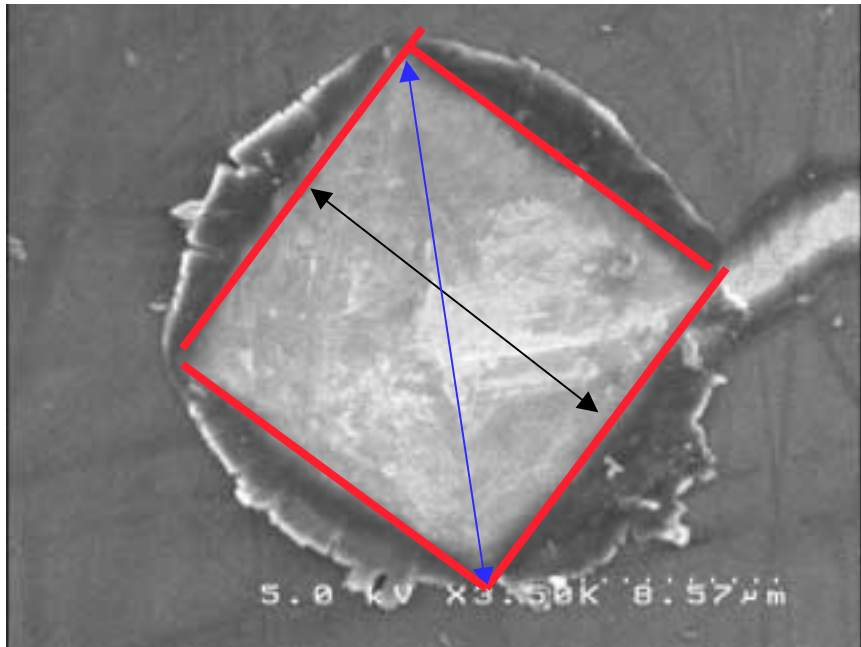
# Vicker Hardness behaviour

## new measurements

Now easier to measure on SEM images (less subjective)

Can measure both side to side and diagonal

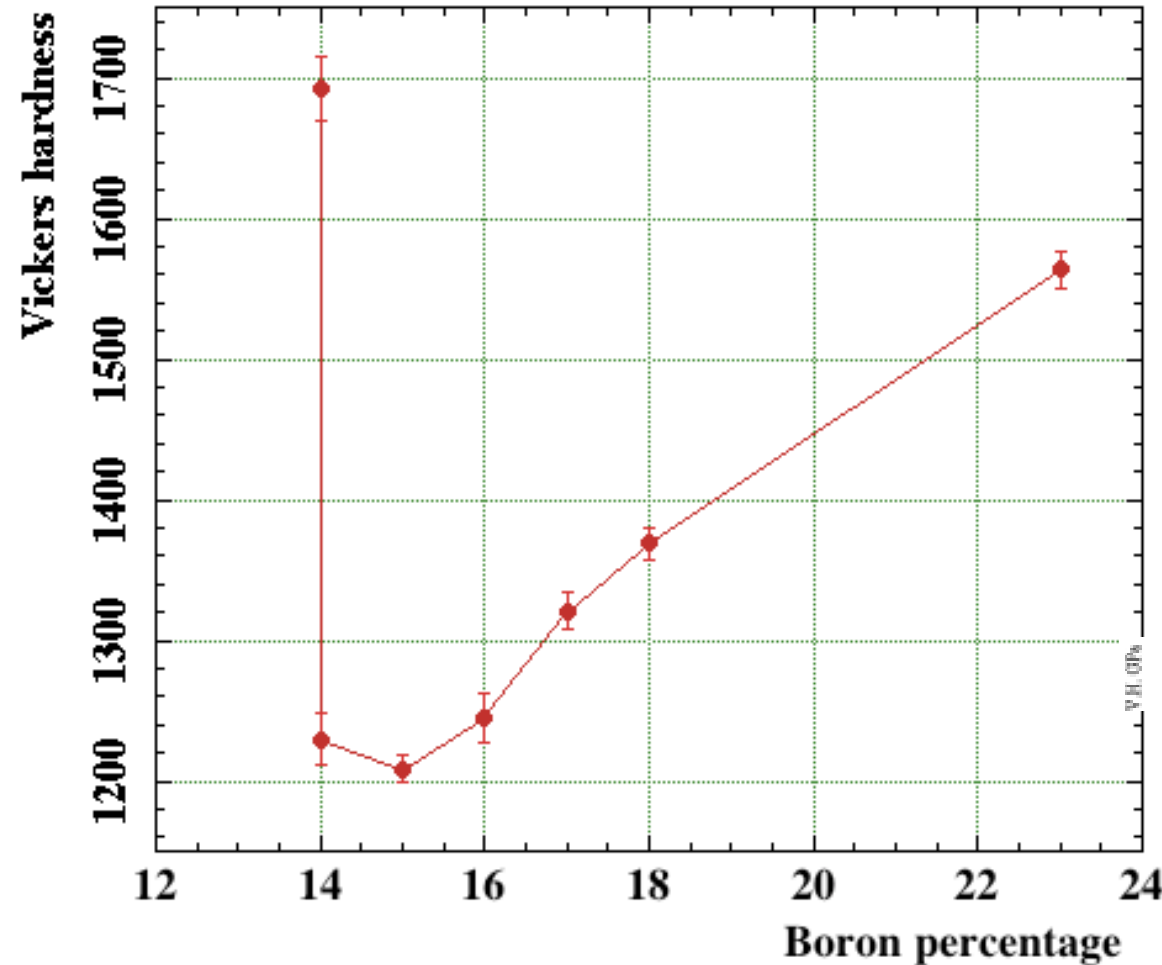
View an plastic outflowing (shear bands) that was previously completely unsuspected and characteristic of different B concentrations



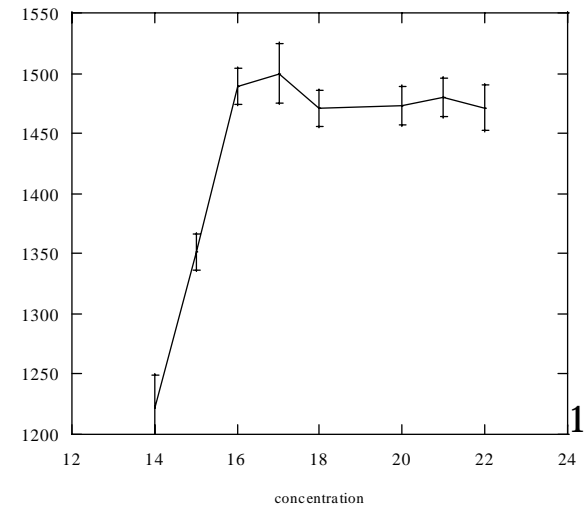
# Vicker Hardness behaviour

## new measurements

MoRuB hardness



Different behavior  
 Dual phase for B14  
 More repetitive  
 Smaller errors



# The braze problem

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- Initially thinking of sputtering because of oxide layer problem
- Found that with aggressive attack and immediate braze coating could avoid the effort (possibly a dead alley)
- Made an oxygen free glove box

LIG



QuickTime™ and a Photo - JPEG decompressor are needed to see this picture.

# AuSn problems

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- Found that despite oxygen free box
- AuSn wets well but peels off
- Checked that AuSnTi wets very well, but also peel off!!!
  
- Lots of wasted time
- But will be faster to reconfigure to AgSnTi

# AgSnTi problems/advantages

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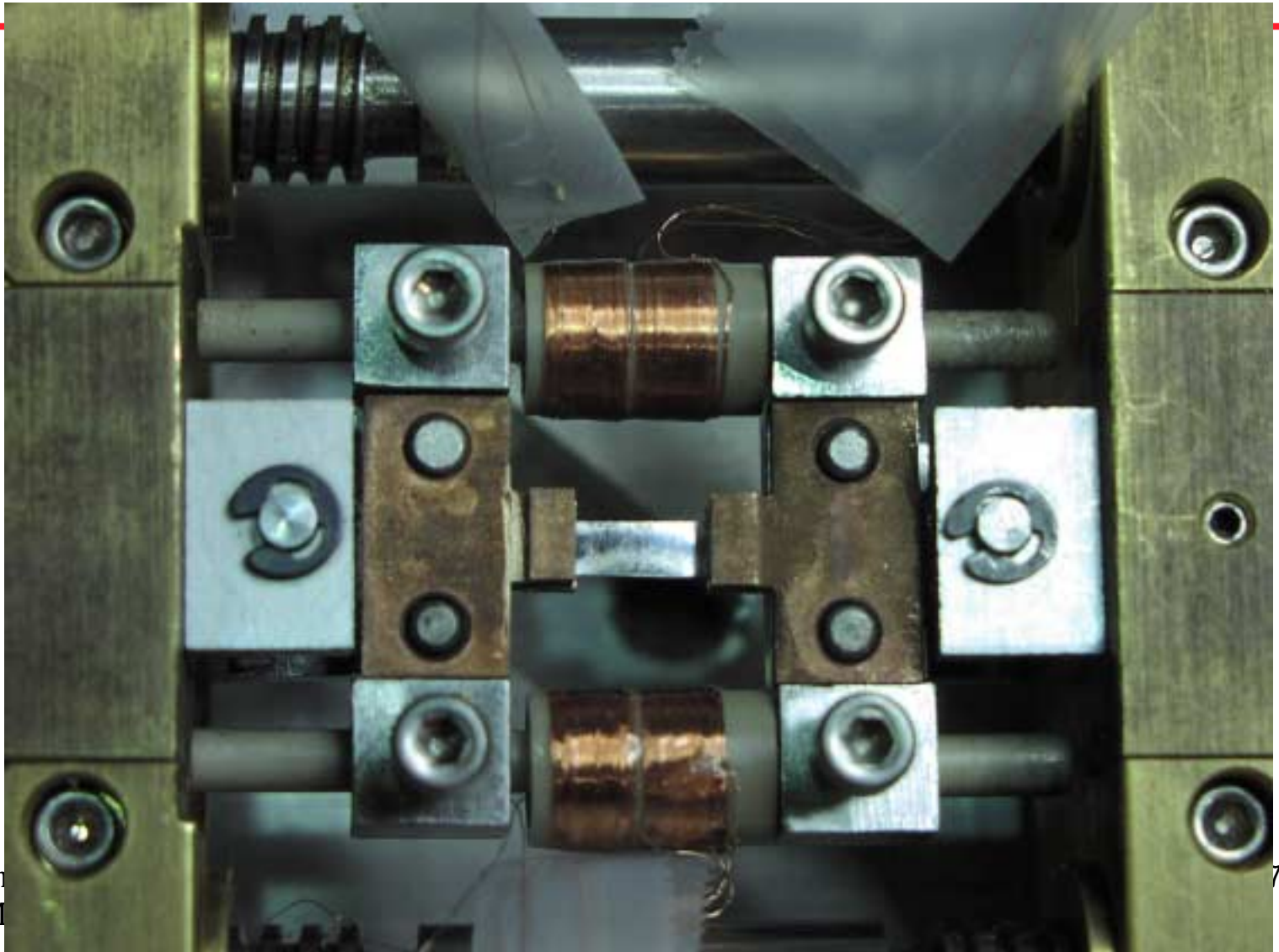
- AgSnTi wets well and STICKS TO IT
- More subject to impurities and pollution
- Softer, will need better control of thickness
  
- If everything fails may need to revert to sputtering

# Stress Strain tests

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- Made tests on MoRuB even if often the braze slipped off cleanly or cracked



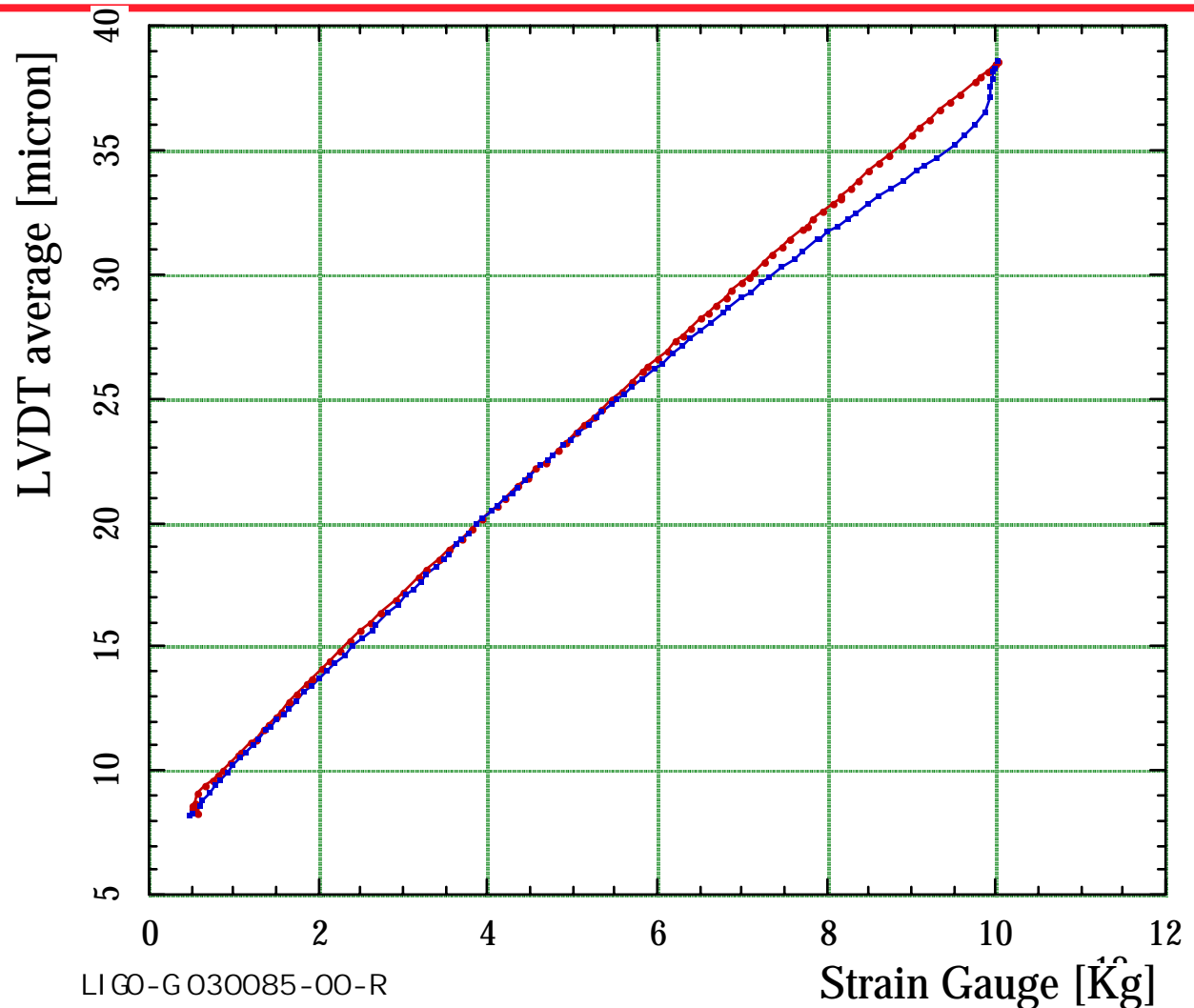




stress strain measurement

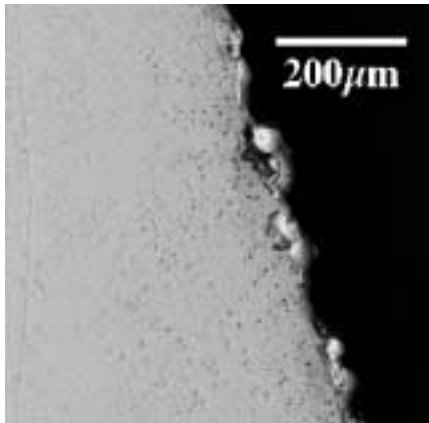
So far achieved  
>2.2 GPa yield  
While expecting  
5.2 GPa

Sample either  
slipping or  
cracking

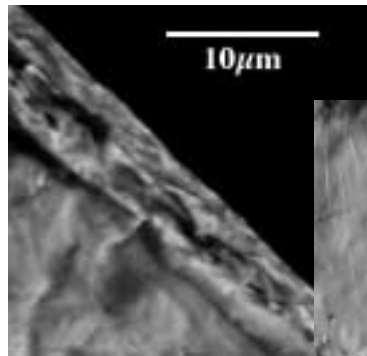


# Why low values for yield point?

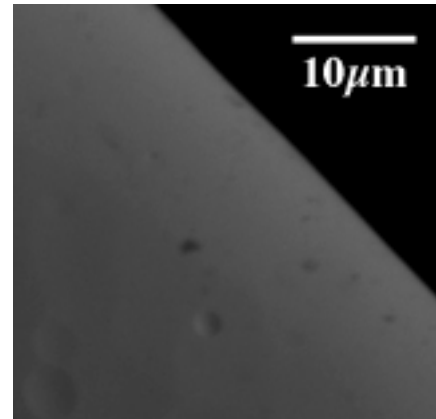
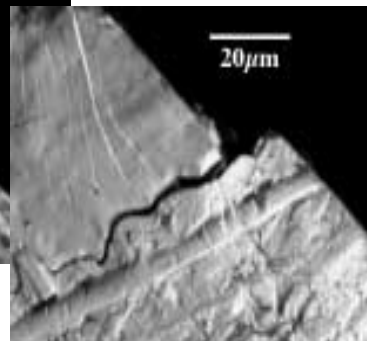
**Nucleation of cracks. To take good measurements we need regular borders without weak point for crack nucleation:**



**EDM Cut**  
Local melting  
Possible formation  
Of crystals on edges



**Scissor cut:**  
Very irregular and unreliable!



**Electropolished cut:**  
The best!

**Nucleation of cracks causes premature failure of the material!**



- Engineering studies  
of suspension structures

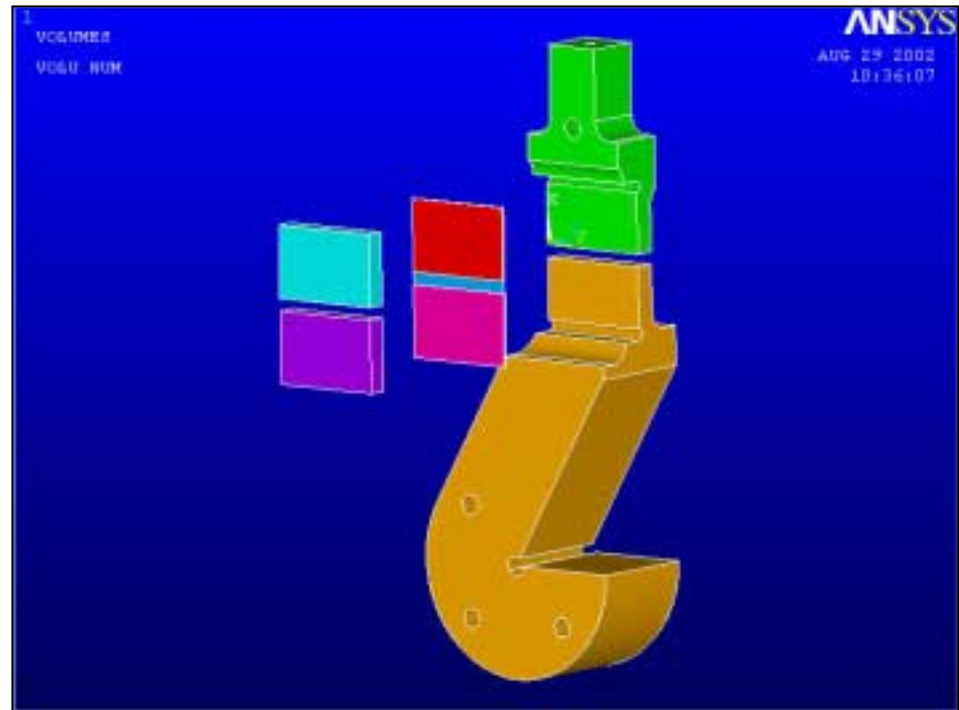
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QuickTime™ and a  
Photo - JPEG decompressor  
are needed to see this picture.

# LIGO Finite Element Analysis of MoRuB flex joint

## Monoflex Assembly:

- Hook
- MoRuB Membrane
- Cavaliers



# Finite Element Analysis of MoRuB membrane

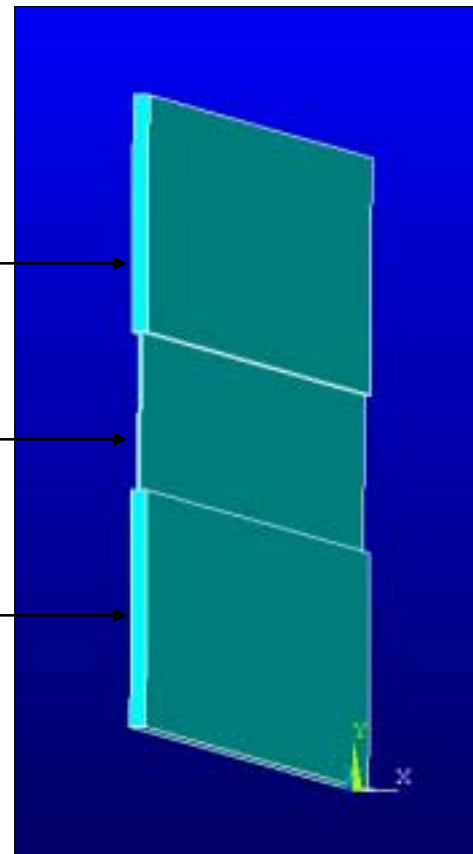
## Initial Membrane Geometry:

- Height: 6mm
- Width: 3mm
- Thickness: 50 $\mu$ m

<10 $\mu$ m

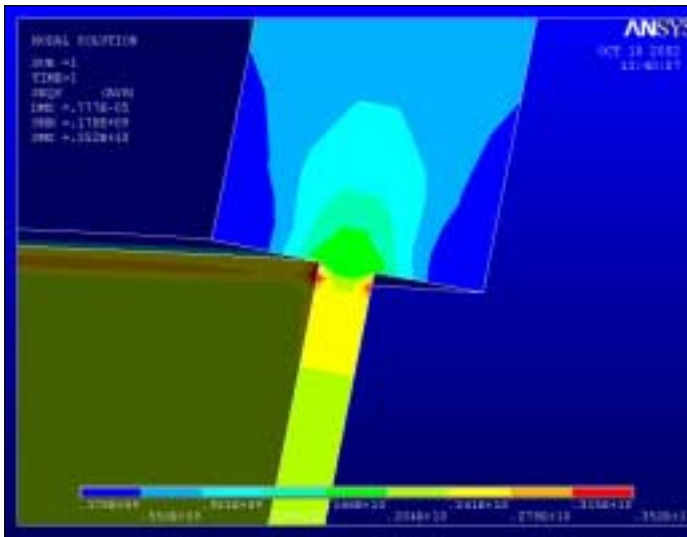
50 $\mu$ m

- With and without fillet radius



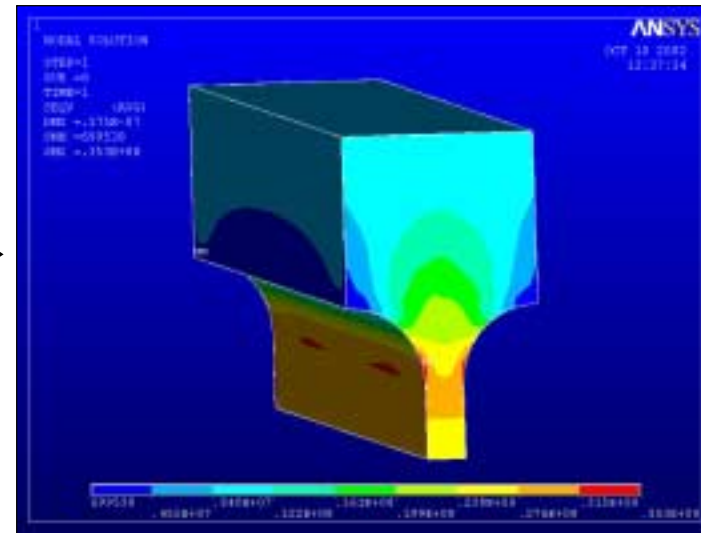
# Filletlet radius effects

In order to minimize the stress concentration in the transition zone a 20 $\mu\text{m}$  fillet radius was chosen.



Sharp transition model

Stress maximum (Von Mises)=3,52e9Pa  
 Min= 2,7e0Pa



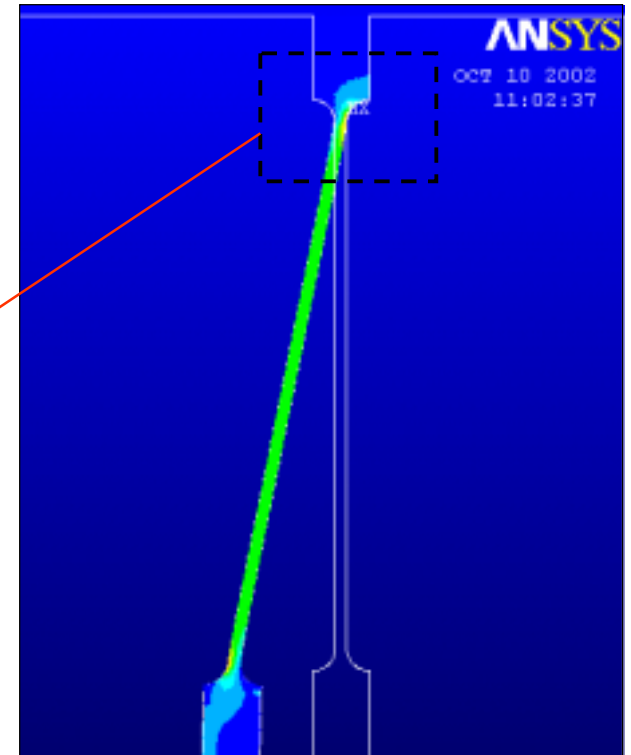
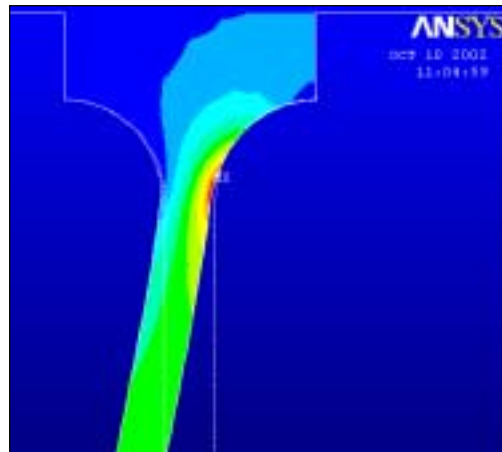
20  $\mu\text{m}$  fillet radius

Stress maximum (Von

# LIGO Finite Element Analysis of MoRuB membrane

## Effective bending length simulation

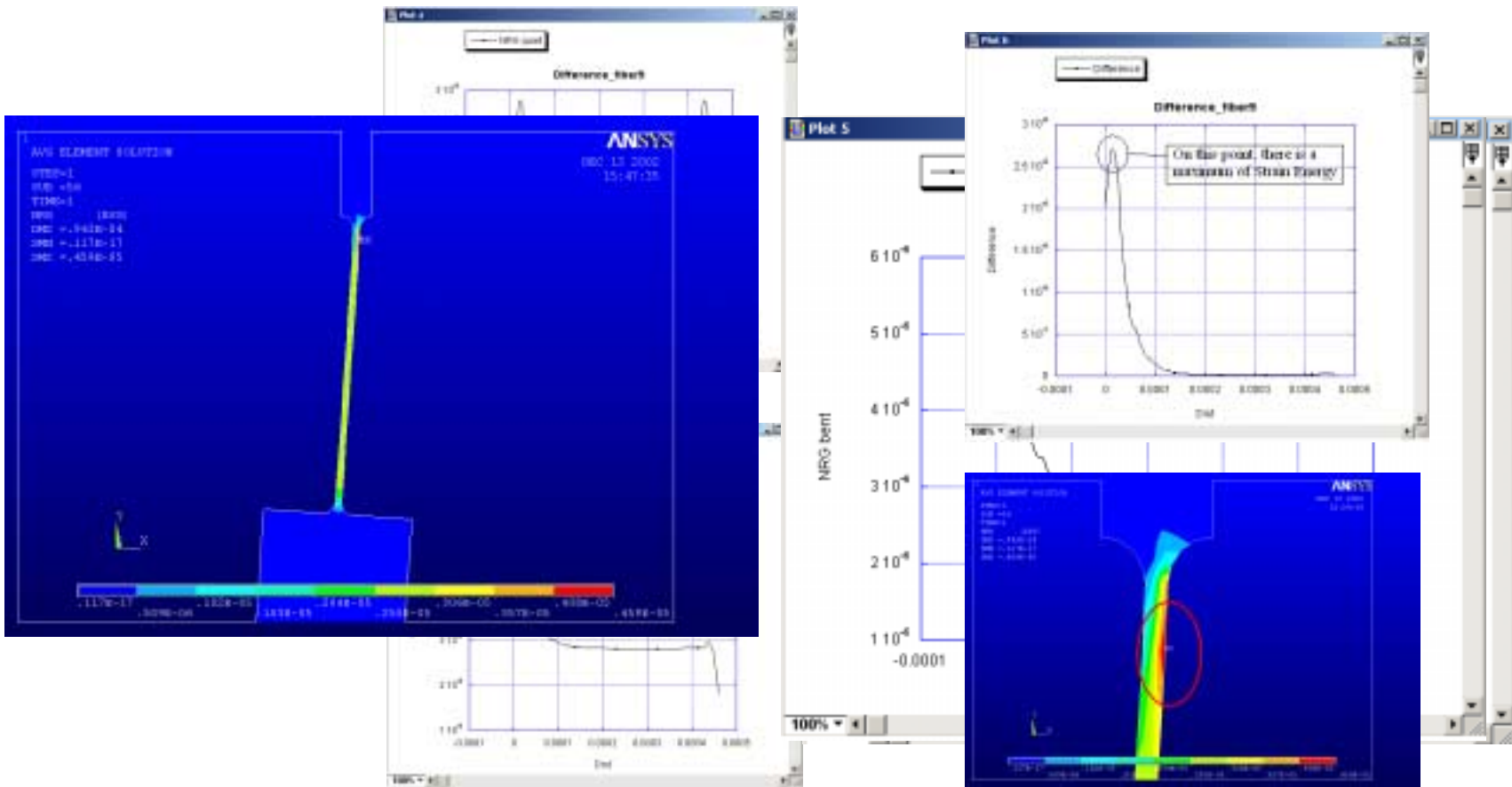
Most oscillation energy stored  
in a length equal to a few thicknesses  
Allowable oscillation of a few mm





# Location of Strain Energy

## The principle



*"Bent" position*

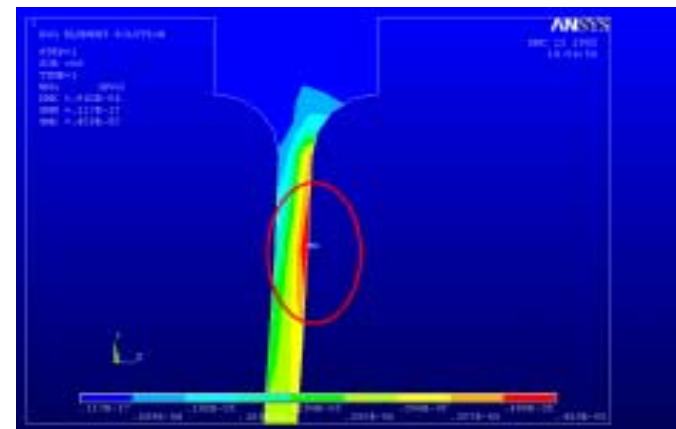
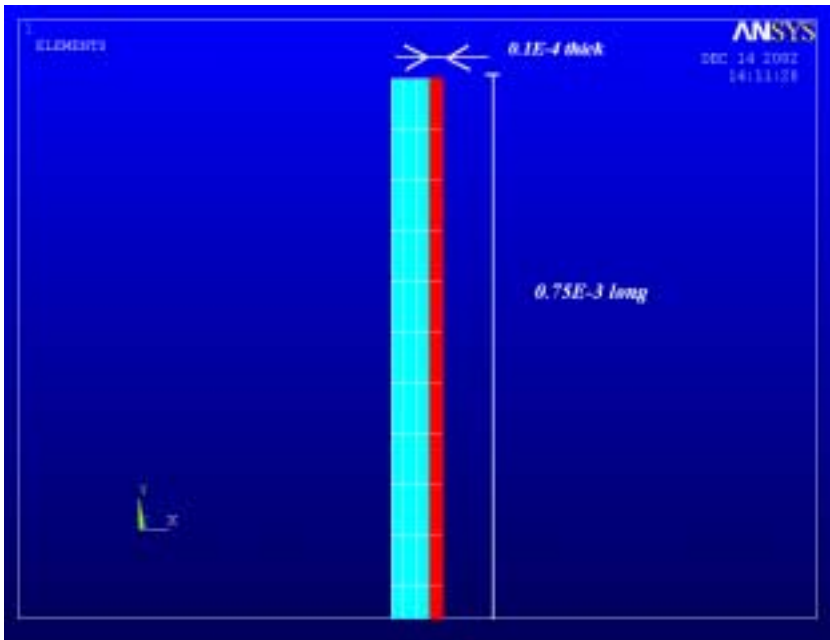
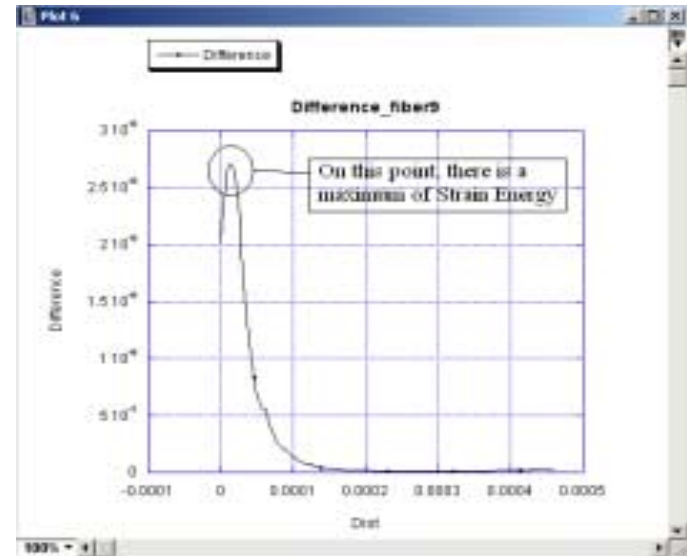
LIGO-G 030085-00-R

# Braze effects

Compare the oscillation strain energy stored in the braze and in the F-joint (Q-normalized)

Effects on overall Q-factor

Thin and wide constrained braze irrelevant



# Conclusions

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- Glassy metals are a promising technology for fully engineer-able mirror suspensions
- Brazeability has still to be fully tested
- Substantially development necessary before being able to consider it a real and advantageous alternative to FS

# Testing jig

**500 Kg mass**  
Supporting  
**10 Kg mirror**

Observe pitch  
mode

