

# “Examination of a Size-Change Test for Photovoltaic Encapsulation Materials”

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**SPIE Optics + Photonics 2012**

**San Diego, CA**

**2012/8/16**

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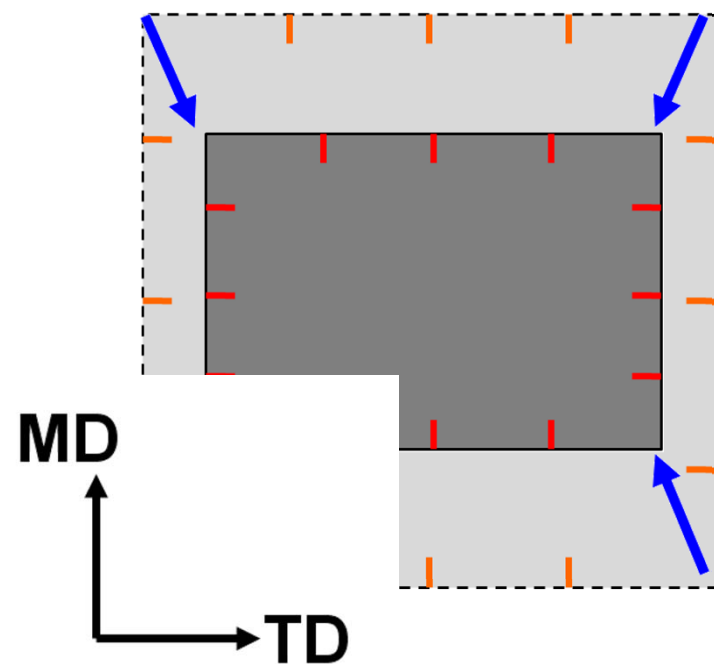
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<sup>8</sup>TÜV Rheinland PTL

**NREL/PR 5200-56320**

# Motivation

- Encapsulation will change size (*e.g.*, shrink) during module processing (lamination)
- Possible consequences for mechanically displaced cells/interconnects/bus-bars: broken solder joints (opens), electrical contact (shunts... cell to cell, ground fault...), cracked cells, delamination, voids in encapsulation
- The long term effects in a field deployed module are unknown
- The encapsulation work-group within IEC TC82 WG2 has proposed a test standard that may be used to assess size change for encapsulation sheet
- Test aids material and module manufacturers in performing material acceptance, process development, design analysis, or failure analysis



*Example demonstrating a size change of  $-45\%$  and  $-20\%$  in the machine extrusion (MD) and transverse (TD) directions*

# Scope and Timeline of the Project

- Measure the maximum representative change in linear dimensions of encapsulation sheet material, resulting from processing during the fabrication of photovoltaic (PV) modules
- A “frictionless” test (between the material and substrate, rendering the maximum size change) is easiest to standardize and interpret
- No existing standard. ISO 11501, ASTM D1204, ASTM D2732 considered

Basis for the test: BP Solar internal test procedure

Task-group formed: Autumn 2010

Discovery experiments and method draft: Spring & summer 2011

Interlaboratory study: Summer and autumn 2011

Method submitted to IEC: Autumn 2011

Revision of draft (from interlaboratory study & IEC vote): 2012

Revised method submitted to IEC: Autumn 2012 or spring 2013

# Details of the Proposed Test Method

- 100 mm x 100 mm specimens: ( $\geq 6$ ), cut from  $\geq 2$  rolls, MD and TD marked, not preconditioned (test promptly)

## Test Procedure:

1. Place Al foil (heat spreader) on hot plate (now a circulating oven)
2. Add 2-4 mm thick layer of sand on Al foil  
weight of sand improves thermal contact of foil  
low friction to standardize the measurement & its interpretation
3. Equilibrate to the maximum processing temperature
4. Measure & record specimen initial dimensions (5 each for MD, TD)
5. Place specimen on sand for 5 minutes
6. Remove, cool then measure specimen final dimensions

7. Calculate size change: 
$$\Delta L = 100 \cdot \frac{L_f - L_i}{L_i}$$

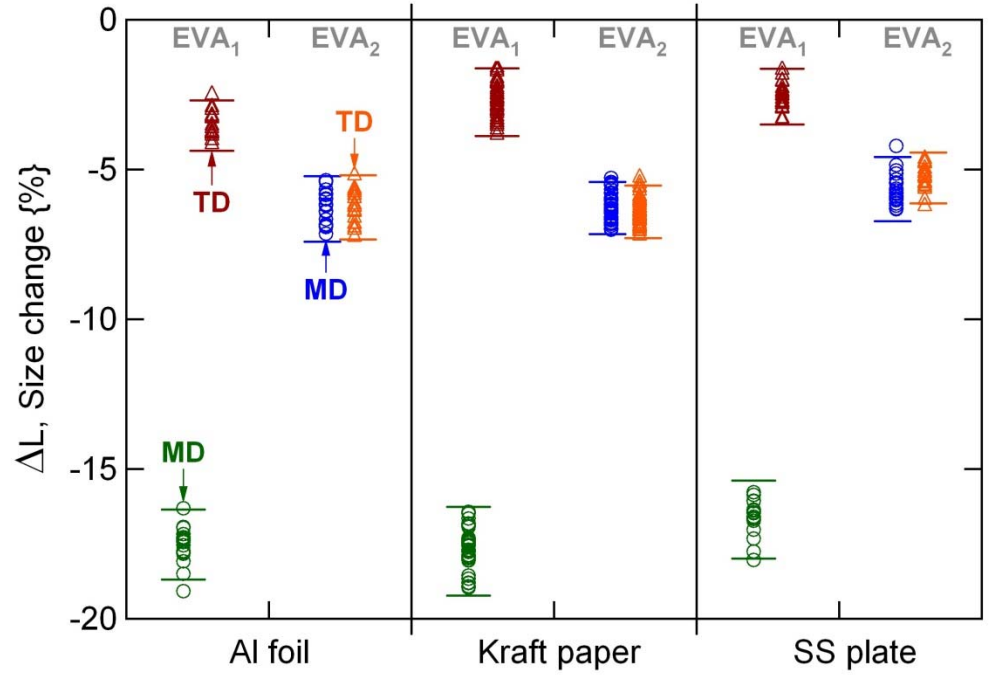
(maximum and difference; average and standard deviation)

# Different “Substrates” Yielded Comparable Shrinkage

- Early work explored talc powder on a glass carrier
- Curvature of glass  $\Rightarrow$  localized thermal contact  $\Rightarrow$  temperature heterogeneity
- Talc is not heavy. Kaolin used in ISO 11501

- Discovery experiment explored sand/carrier combinations for 2 EVA's

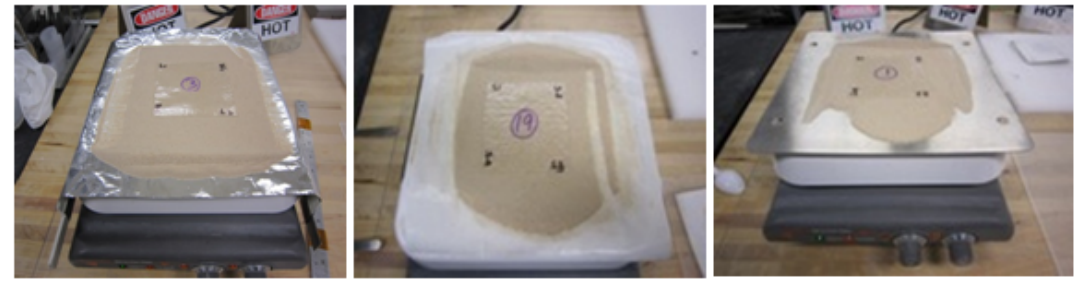
- No significant ( $2\sigma$ ) difference observed. Al chosen.



*Comparison (data and images) of carrier/sand for 2 EVA's (unbalanced and balanced).*

“Kraft paper “= release liner paper

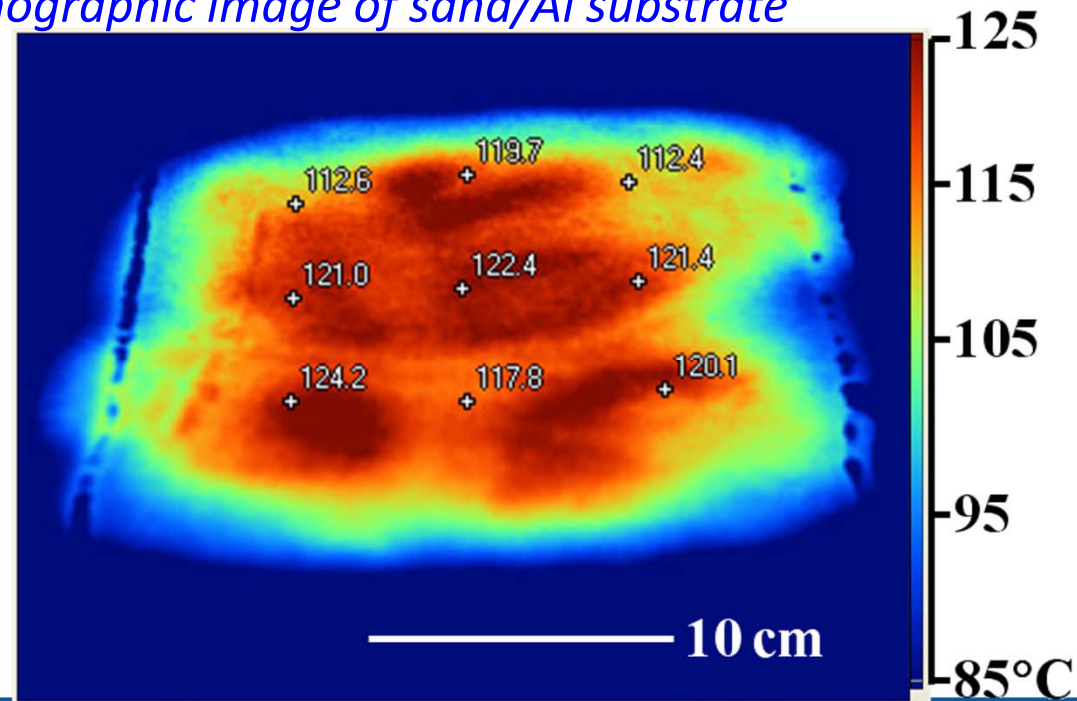
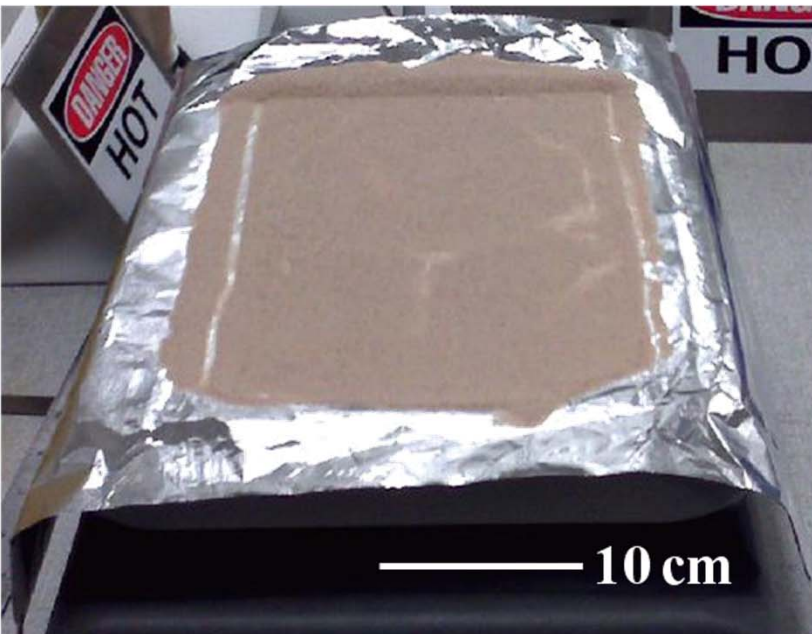
**Aluminum Foil      Kraft Paper      Stainless Steel Plate**



# Quantifying the Temperature Uniformity of Sand

- Sand (unlike Al) is a high  $\epsilon$  material, readily enabling thermography
- A 4-8°C ( $2\sigma$ ) T range was observed for well manicured sand
- Most heterogeneous at thin regions or for partially raked sand
- Circulating oven can improve temperature stability and uniformity:
  - no temperature gradient through the sand, no radiative heat transfer,
  - greater thermal capacitance, better recovery time, safety

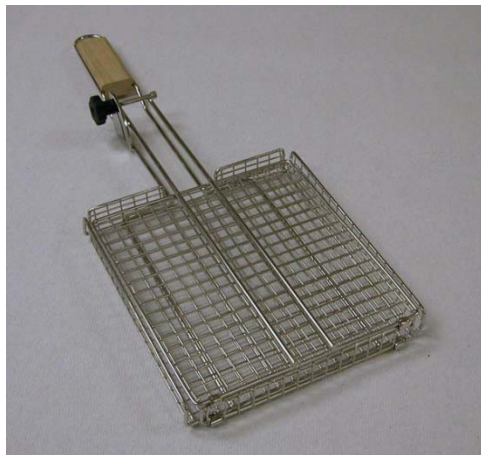
*Optical and corresponding thermographic image of sand/Al substrate*



# The Possibility of a Liquid “Substrate” Seems Unlikely

Scenario:

- A liquid-based test was identified (as in ASTM D2732) from the IEC vote
- Some voters advocated the use of water (@ 80°C) to evaluate EVA



*wire tray ([www.eysters.com](http://www.eysters.com))*



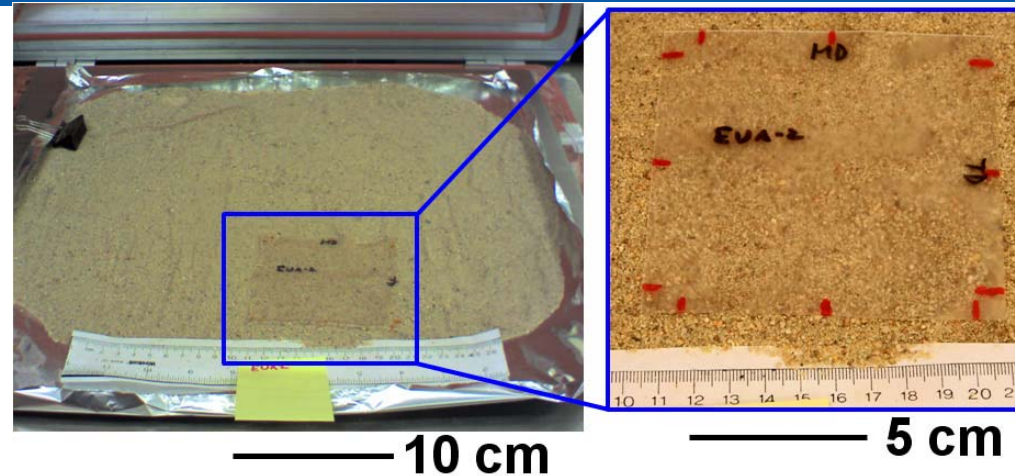
*wire basket ([www.sockmete.info](http://www.sockmete.info))*

Difficulties:

- 80°C not expected to cure EVA.  $\Delta L$  from melt transition only.
- The standard is intended to rapidly test all encapsulation materials (including those that are processed at >100 °C)
- Many new encapsulation materials do not cross-link and are processed near/above their melting temperatures
- How to handle molten materials without introducing shape change?

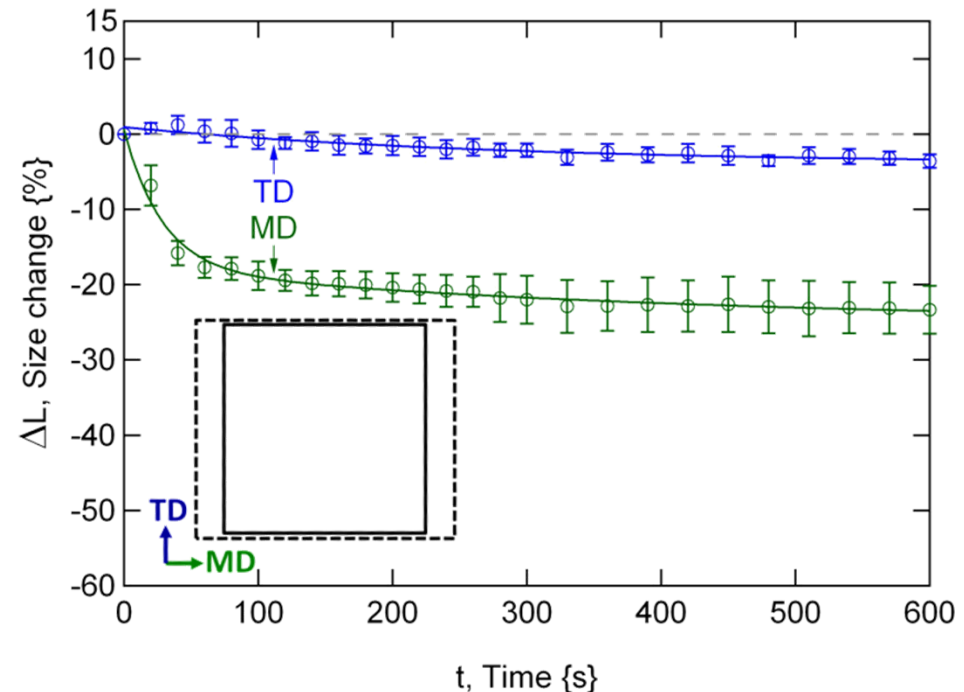
# Experiments Confirm the Test Duration for EVA

- Photographs taken every 20s for specimens marked at middle and near the corners
- Size change can be determined optically ( $\sim\pm 1\%$ ) using the scale in the image



“Hot plate” (vacuum laminator) and specimen setup for the time characterization

- Experiment temperature =  $132^{\circ}\text{C}$
- Negative  $\Delta L$  = shrinking
- The initial (dashed) and final (solid) profiles are shown (scaled) in the figure inset
- EVA: most activity within 1<sup>st</sup> two minutes (EVA cross-links)

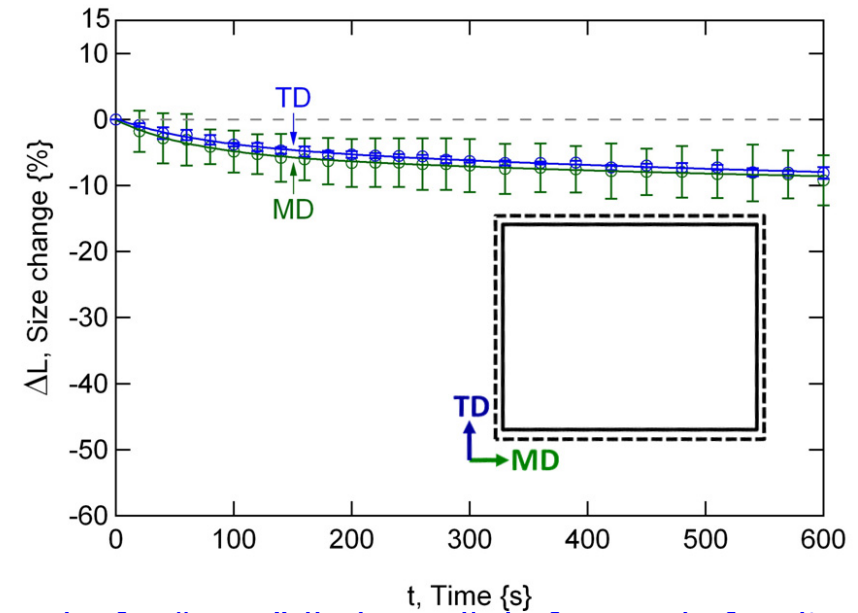


Results for “EVA<sub>1</sub>” (unbalanced), before and after (inset). Error bars shown for max and min measurements.

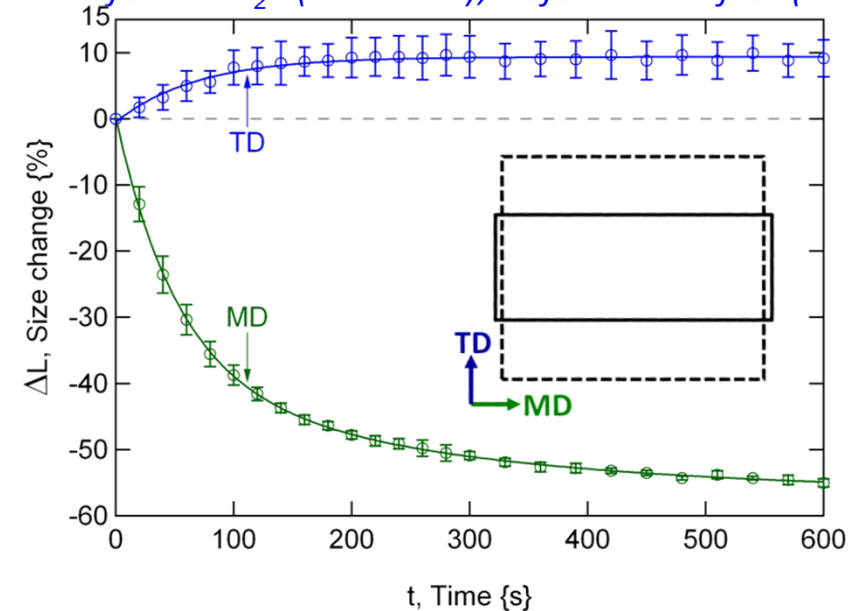


# Experiments Confirm the Test Duration for Other Encapsulation

- The thermoplastics do not cure, but demonstrate most size change within 5 minutes
- Some materials tested at 165°C
- Examples:
  - balanced EVA (10% → minimal size change)
  - TPO (55% → substantial size change)
- Some materials (e.g., TPO, PVB, and ionomer thermoplastics) shrink in one direction and expand in the other!
- Some materials not optimized to reduce size change, as vendors are likely unaware of the issue
- The implications for the stress in a module are unclear (try FEA) but may become more significant with time



Results for "EVA<sub>2</sub>" (balanced), before and after (inset)



Results for TPO, before and after (inset)

# A Minor Specimen Size-Effect is Evident

## Experiment:

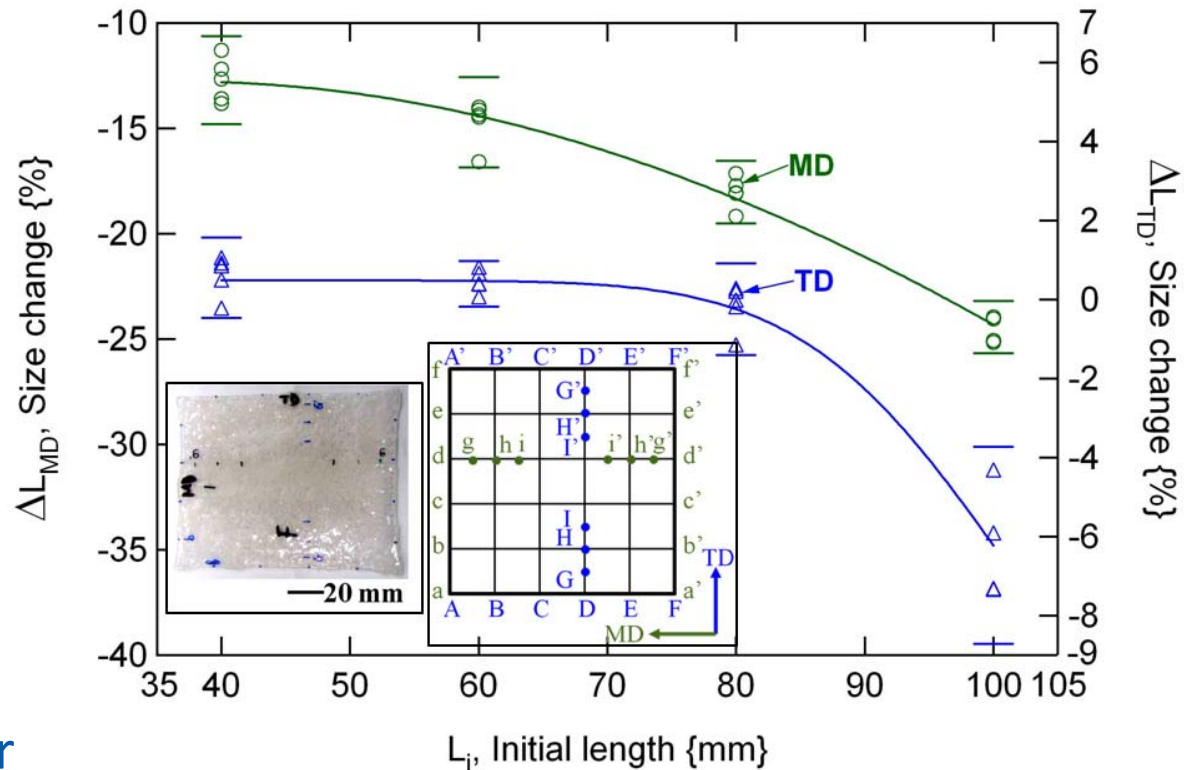
- Is there a size-effect?
- Obtain measurements from within and up to the specimen edges
- $L_i = 40, 60, 80, 100$  mm

## Results:

- $\Delta L$  at edge for EVA's (as in TD for EVA<sub>1</sub>)
- Monotonic trend for "isotropic" PVB, TPO, ionomer (like MD for EVA<sub>1</sub>)

## Implications:

- Possible causes: friction (from sand), stretching during cutting, uneven & rapid cooling, heterogeneous stress



Size-specific results for "EVA<sub>1</sub>" (unbalanced).

(a) Final photograph of one of the specimens.

(b) Sign convention and coordinate system used.

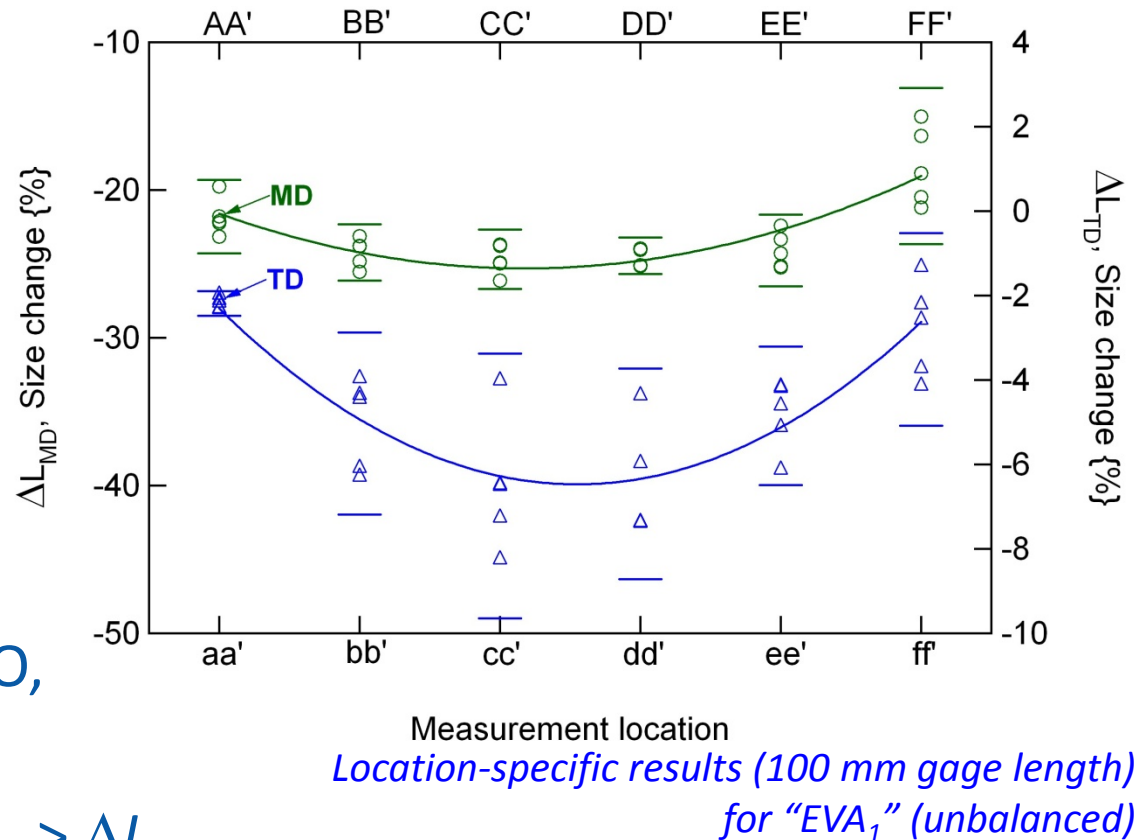
# A Minor Edge-Effect is Evident

## Experiment:

- Intentionally measure at locations along specimen edges, including the corners
- A minor effect (few % $\Delta L$ ) is evident in all specimens

## Results:

- Similar behavior for EVA, TPO, ionomer:  $\Delta L_{DD'} > \Delta L_{AA'}$
- Opposite trend for PVB:  $\Delta L_{AA'} > \Delta L_{DD'}$

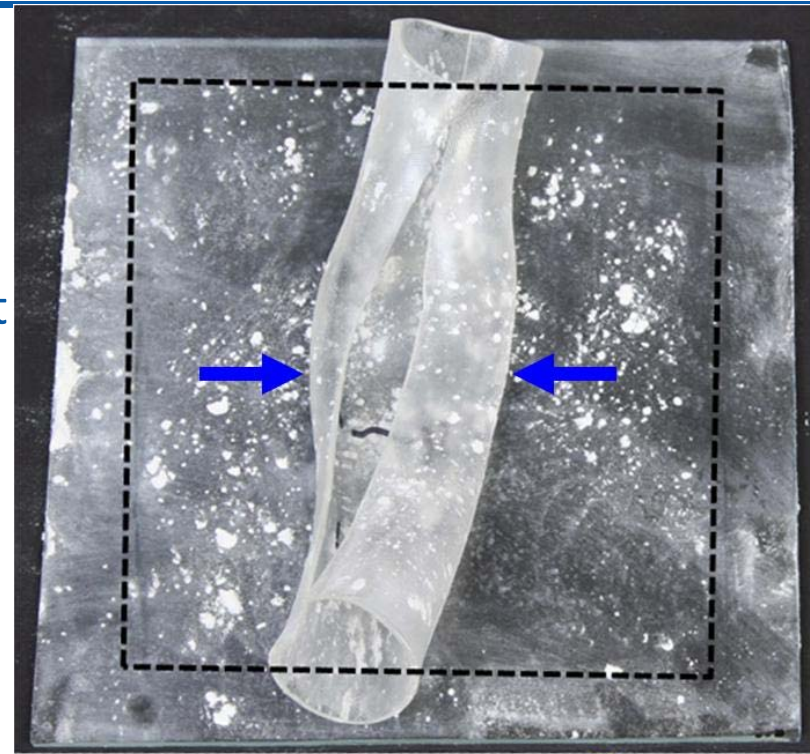
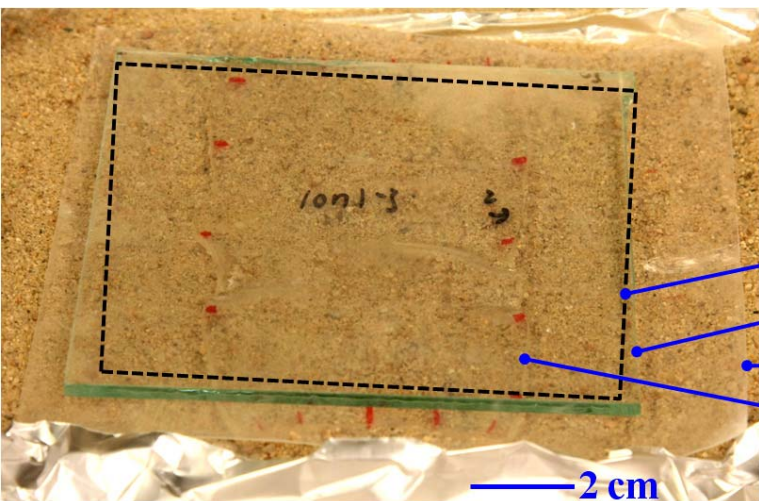


## Implications:

- Specify the # and location of measurement sites
- Measure middle and  $\geq 1$ cm from the corners, using an odd # sites)
- Sample  $\geq 200$ mm (location) from the edge of a roll

# How to Treat Out of Plane Curvature?

- Early generation ionomer product:  
 $\Delta L_{MD} \leq -50\%$ ,  $\Delta L_{TD} \geq 15\%$ , significant curvature
- $\Delta L$  could probably be significantly improved
- Not practical to uncurl and measure @ end of test
- For in-plane result, one could cover with Teflon FEP sheet /weight (e.g., glass)



— 2 cm  
*Image of final shape of ionomer (arrows at edges), with outline (dashes) of original shape*

*Image of final shape of ionomer/FEP/glass*

- This solution does, however, affect the result (magnitude and material profile)
- Are there better practices? Note: glass weight often not “required”

# Details of the Interlaboratory Study

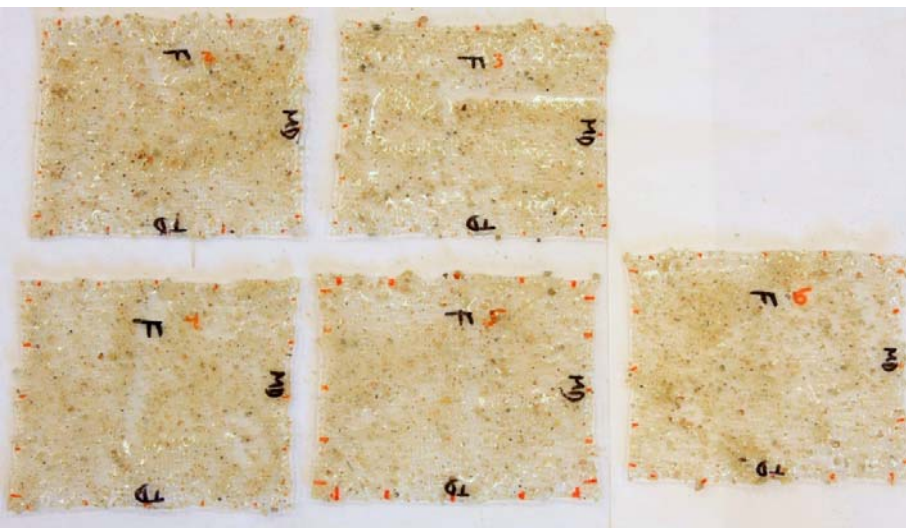
- Five materials were circulated:

EVA<sub>1</sub> (unbalanced;  $T_{\text{set}}=132^{\circ}\text{C}$ ;  $T_{\text{m}}=55^{\circ}\text{C}$ ), EVA<sub>2</sub> (balanced),

←thermosets; thermoplastics→ TPO ( $T_{\text{set}}=140^{\circ}\text{C}$ ;  $T_{\text{m}}=60^{\circ}\text{C}$ ),

PVB ( $T_{\text{set}}=160^{\circ}\text{C}$ ;  $T_{\text{g}}=15^{\circ}\text{C}$ ), ionomer ( $T_{\text{set}}=165^{\circ}\text{C}$ ;  $T_{\text{g}}=86^{\circ}\text{C}$ )

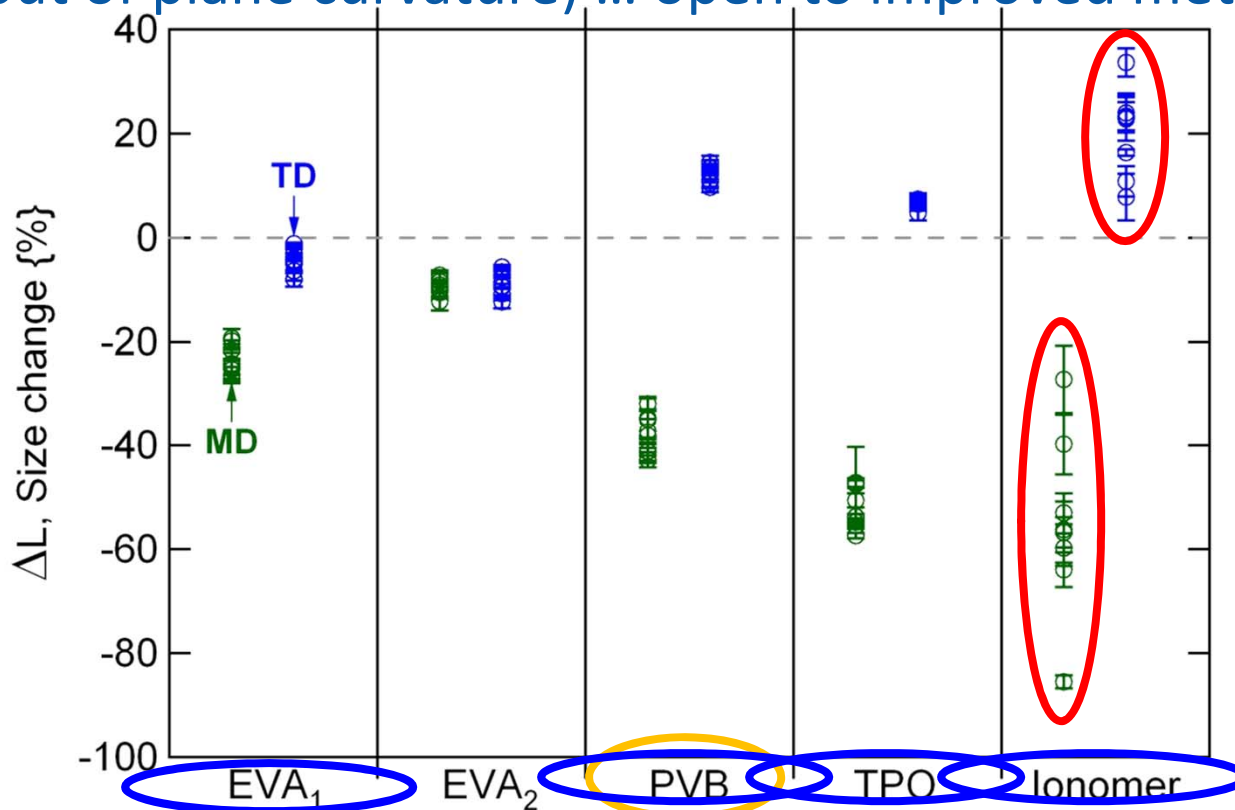
- $\Delta L$  measurements for MD, TD according to the draft procedure
- Tests were performed using a hot-plate or oven with Al foil
- Unspecified sand substrate (now ASTM C778)



*EVA<sub>1</sub> (unbalanced) specimens after the test*

# Results of the Interlaboratory Study

- Most materials (except PVB) were examined in the melt state
- $\Delta L_{MD} > \Delta L_{TD}$  for EVA<sub>1</sub>, PVB, TPO, ionomer
- Results are reproducible between participating laboratories (within  $\pm 5\%$  absolute [from  $L_i$ ], up to 40% relative [from  $\Delta L$ ])
- The ionomer was not very repeatable between labs (out of plane curvature) ... open to improved method for this issue!



*Box plot of average and st dev of size-change from the interlaboratory study*

# Summary

- Proposed test standard to evaluate the maximum change in linear dimensions of sheet encapsulation products resulting from their thermal processing. Discovery and interlaboratory studies performed.

Sand substrate, aluminum carrier:

- Reduce friction (maximum size change) standardizing the test
- Sand can be used at a wide range of test temperatures
- Specify to use circulating oven
- We anticipate a 5°C (2 $\sigma$ ) range within the oven

Related details:

- Verified 5 minute duration for the test
- Minor size-, edge-effects  $\Rightarrow$  specify size, measurement locations
- Difficult to reduce effects of out-of-plane curvature

Interlaboratory study:

- Substantial size change (>10%) observed for several materials
- Often observed shrinking in MD, expansion in TD
- Results reproducible within  $\pm 5\%$  absolute size-change

# Acknowledgments

- NREL: Dr. Michael Kempe, Dr. Sarah Kurtz, Dr. John Pern, Steve Glick



This work was supported by the U.S. Department of Energy under Contract No. DE-AC36-08GO28308 with the National Renewable Energy Laboratory.

See also the manuscript: “Examination of a Size-Change Test for Photovoltaic Encapsulation Materials”, Proc. SPIE 2012, 8472-29.