

# Effect of CMP on abrasives and their surface characteristics

---

S.V. Babu

Center for Advanced Materials Processing  
Clarkson University



# Outline

- Introduction
- Common CMP slurry abrasives and additives
- Effect of additives and polishing on abrasives
- Some aspects of III-V polishing
- Contribution of pad to slurry waste
- Summary and Conclusions
- Acknowledgements

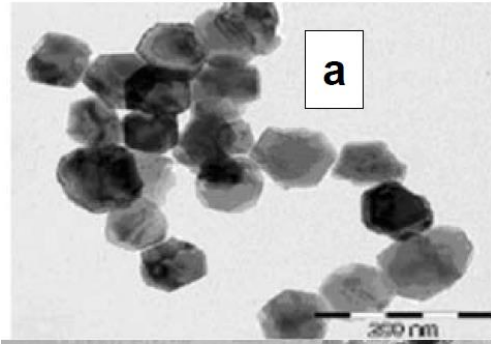
# Common slurry components: BEO

Substrate	Slurry Components	Species present post-polish
<ul style="list-style-type: none"> <li>• Cu</li> <li>• Cu/Ta/TaN</li> <li>• Cu/Ti/Co</li> <li>• Cu/Ti/Ru</li> <li>• Cu/Mn</li> <li>• W</li> <li>• SiCOH</li> </ul>	<ul style="list-style-type: none"> <li>• Silica/Alumina</li> <li>• pH adjusting agents such as HNO<sub>3</sub>, KOH, NH<sub>4</sub>OH, etc. and buffers</li> <li>• Oxidizer (H<sub>2</sub>O<sub>2</sub>, Ferric nitrate, KIO<sub>4</sub>, KMnO<sub>4</sub> etc.)</li> <li>• Complexing agents (amino acids such as glycine, carboxylic acids such as citric acid)</li> <li>• Corrosion inhibitors ( BTA and its derivatives)</li> <li>• Surfactants</li> </ul>	<ul style="list-style-type: none"> <li>• OH<sup>-</sup>, surfactants/ adsorbed metal ions/ metal oxides and pad debris</li> <li>• dissolved, suspended and settled fine particles and dissolved metals</li> <li>• Unreacted oxidizers, complexing agents, inorganic ions, etc.</li> </ul>

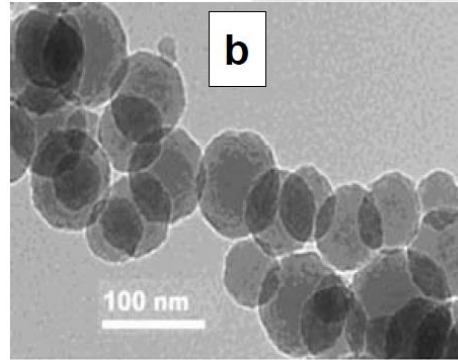
# Common slurry components: FEOL

Substrate	Slurry Components	Species present post-polish
<ul style="list-style-type: none"><li>• <math>\text{SiO}_2/\text{Si}_3\text{N}_4</math></li><li>• Poly-Si</li><li>• W, Al</li><li>• Ge</li><li>• GaAs</li><li>• InP</li><li>• InGaAs</li><li>• InAlAs</li></ul>	<ul style="list-style-type: none"><li>• Silica, Ceria</li><li>• pH adjusting agents such as <math>\text{HNO}_3</math>, <math>\text{KOH}</math>, <math>\text{NH}_4\text{OH}</math>, etc. and buffers</li><li>• Oxidizer (<math>\text{H}_2\text{O}_2</math>, etc.)</li><li>• Surface modifying agents</li></ul>	<ul style="list-style-type: none"><li>• Arsenic rich waste</li><li>• Dissolved <math>\text{AsH}_3</math>, <math>\text{PH}_3</math> and <math>\text{NH}_3</math></li><li>• Adsorbed metal ions</li><li>• Unreacted Oxidizing agents</li><li>• Inorganic ions</li><li>• Unreacted complexing agents</li></ul>

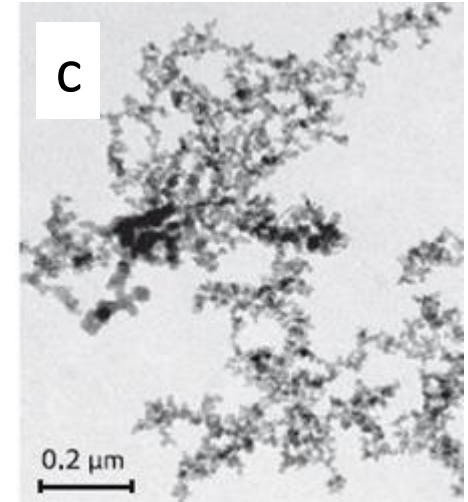
# Common abrasives used in CMP and their characteristics



(a) Rhodia Ceria  
( $d_m$  -60nm)



(b) Nexsil colloidal Silica  
( $d_m$ -50 nm)

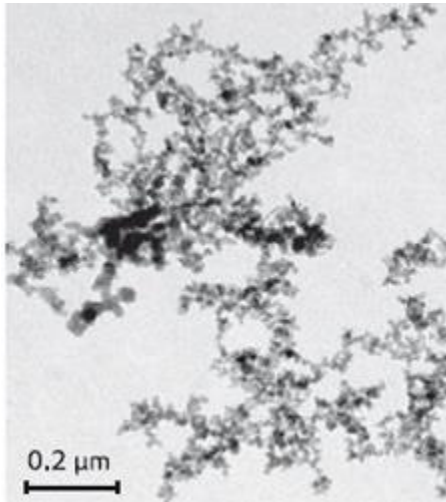


(c) Aerosil Fumed Silica  
( $d_m$ - 130 nm)

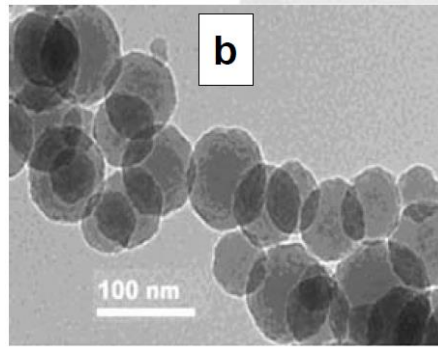
	<b>Bulk density (g/cm<sup>3</sup>)</b>	<b>Mohs Hardness</b>	<b>IEP</b>	<b>Possible species on surface</b>
Silica	2.65	6-7	~2	$\text{M-OH}_2^+ \longrightarrow \text{M-OH} + \text{H}^+$ $\text{M-OH} \longrightarrow \text{M-O}^- + \text{H}^+$ <p style="text-align: center;"><b>M= Metal</b></p>
Ceria	7.65	6	6-8	
Alumina	3.95	9.0	9	

- *H. Bergna et al., Colloidal Silica Fundamentals and Applications, Taylor and Francis, 2006*
- George V. Franks, and Yang Gan *J. Am. Ceram. Soc.*, 90 [11] 3373–3388 (2007).

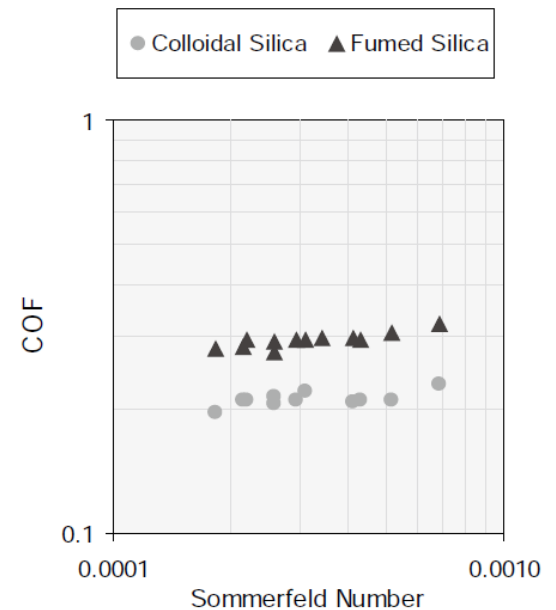
# Differences between colloidal and fumed Silica



Aerosil Fumed silica



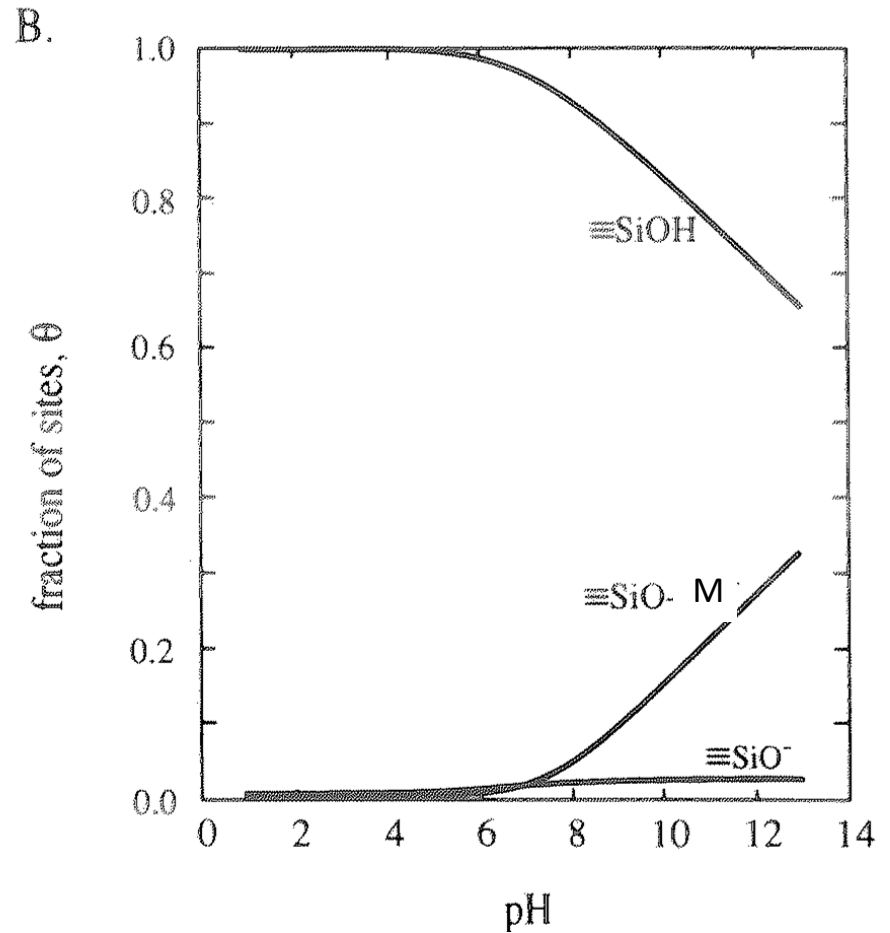
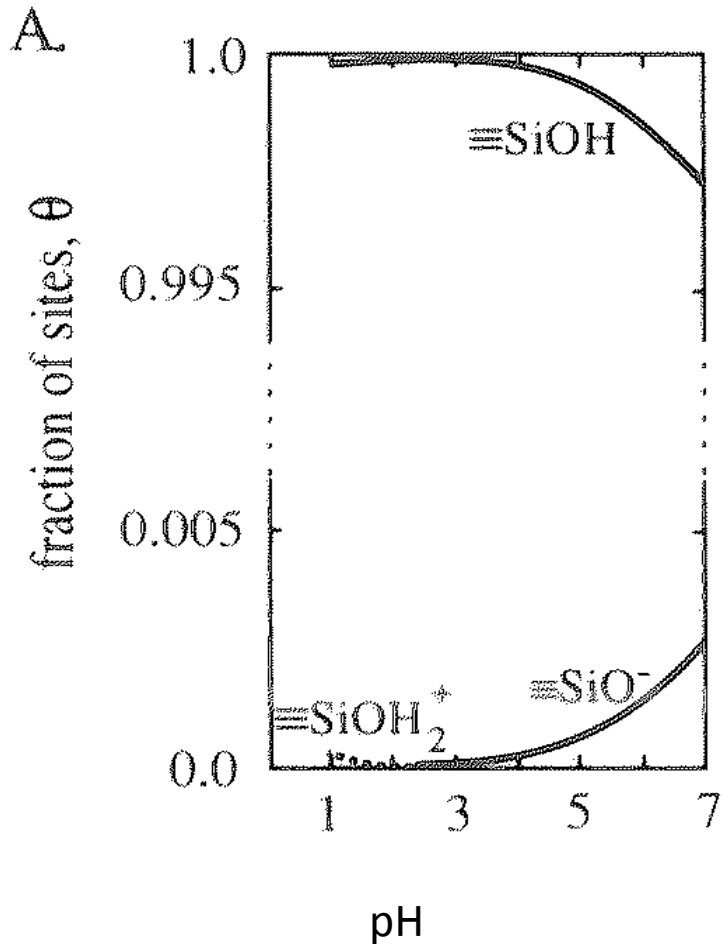
Nexsil 85A Colloidal silica



Fumed Silica	Colloidal Silica
Possible <b>Cl</b> contamination due to manufacture from chloro-silane (<200 ppm)	Possible <b>Na</b> contamination due the to manufacture from sodium silicate
Large hard and soft aggregates Generally used at high pH -> High Concentration of dissolved silica	Surfactants to maintain colloidal stability

- <http://www.aerosil.com/product/aerosil/en/industries/papers/pages/default.aspx>
- Z. Li, K. Ina, P. Lefevre, I. Koshiyama, and A. Philipossian. Journal of The Electrochemical Society, 152(4)G299-G304~2005.

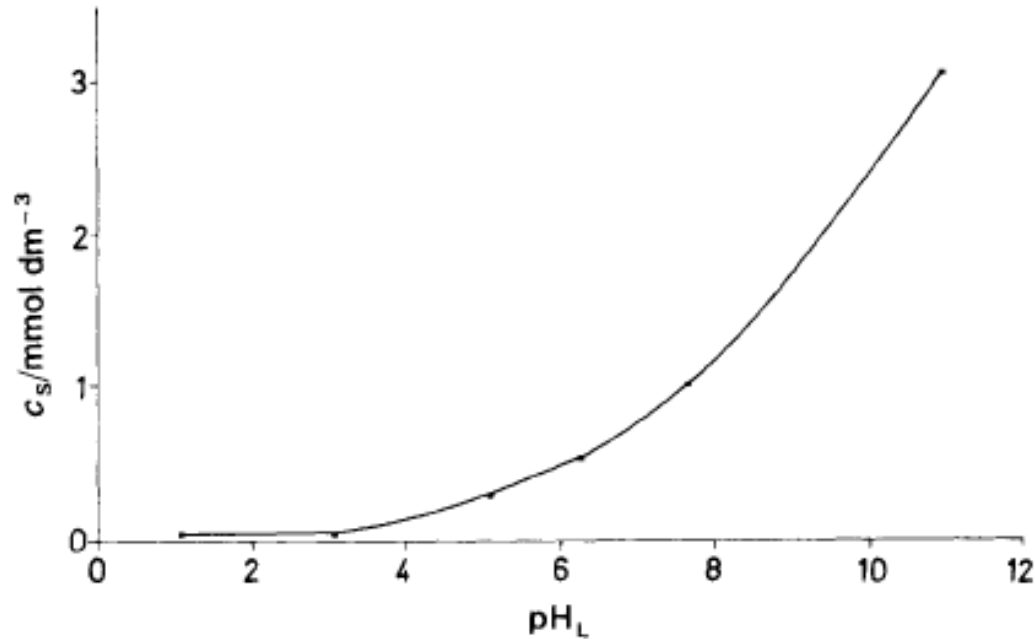
# Silica Speciation



M= Cation from the pH adjusting agent ( $\text{K}^+$ ,  $\text{NH}_4^+$ )

- Dove and Elston. *Geochimica et Cosmochimica Acta*. Volume 56, Issue 12. December 1992, Pages 4147–4156.

# Solubility of Silica as a f(pH)



## Saturation concentration of silica, ( $C_s$ ) as a function of pH

Silica can dissolve as a silicate as shown below:



Can form insoluble species quickly when they react with metal cations ( $\text{Ca}^{+2}$ ,  $\text{Zn}^{+2}$  and other heavy metal impurities)

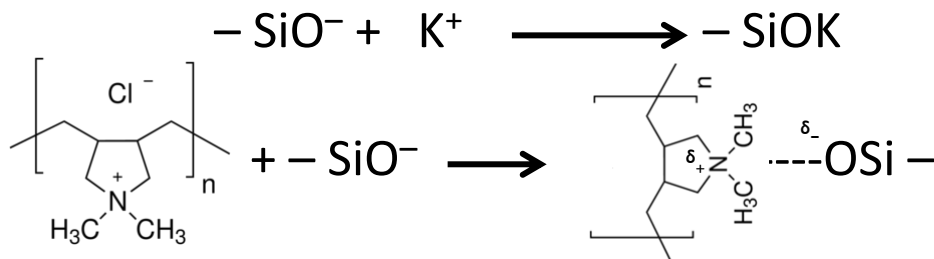
- Wolfram Vogelsberger,\* Andreas Seidel and Georg Rudakoff. CHEM. SOC. FARADAY TRANS., 1992, 88(3), 473-476



# Possible interactions and metal impurities on silica surface

## Electrostatic Interactions :

Adsorption of cationic surfactants/stabilizers and metal ions.



## Acid-Base Reactions:

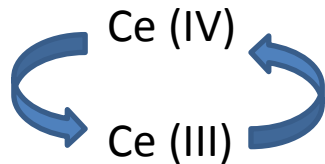


	Metal	Conc. (ppb)
1	Al	50
2	Ca	110
3	Cr	22
4	Cu	10
5	Fe	85
6	Mg	20
7	Ni	11
8	K	50
9	Na	20
10	Zn	11

Trace Metal Contamination in **fumed** silica slurries measured with ICP-AES

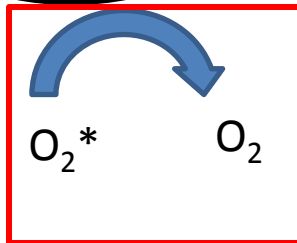
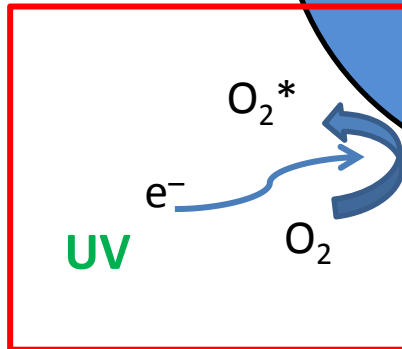
# Possible interactions on ceria surface

## Acid-Base Reactions:

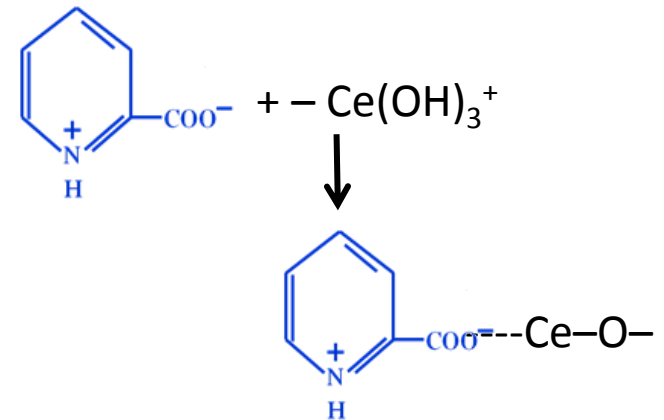


Excellent catalyst for many reactions

Ceria



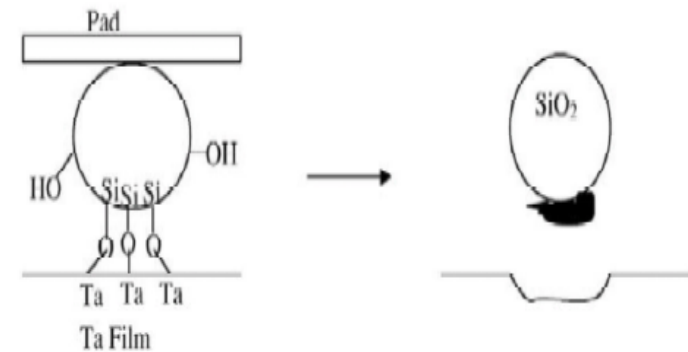
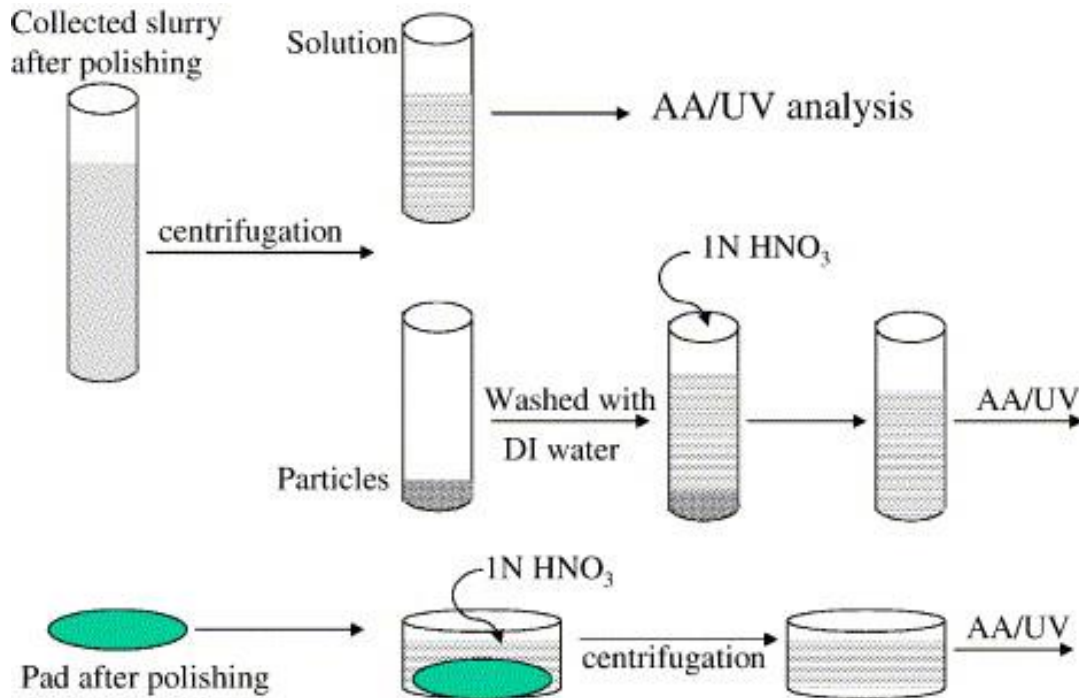
## Electrostatic:



Switchable oxidation states: ROS generation/ ROS consumption

# Effects of polishing on the abrasives

# Cu/Ta barrier CMP - Where does the polished metal go?



Using silica to polish Ta ( $Ta_2O_5$ )

- Ying Li, Junzi Zhao, Ping Wu, Yong Lin, S.V. Babu, Yuzhuo Li. *Thin Solid Films*, Volume 497, Issues 1–2, 21 February 2006, Pages 321–328.
- Y Li, M Hariharaputhiran, S.V Babu. *J. Mater. Res.*, 16 (2001), p. 1066
- A. Vijayakumar, T. Du, K.B. Sundaram, V. Desai, *Polishing mechanism of tantalum films by SiO<sub>2</sub> particles*, *Microelectronic Engineering* 70 (2003) 93-101

# Ta CMP results using modified and unmodified silica in water

<i>Location found</i>	<i>Fumed Silica</i>		<i>Modified silica</i>	
	Ta detected (mg)	% Ta detected	Ta detected (mg)	% Ta detected
Solution	0.33	22	< 0.2	100
Abrasive	1.13	<b>75.3</b>	ND	ND
Pad	0.04	2.7	ND	ND
Total detected	1.5	100	< 0.2	100
Ta loss from disk per polish	1.63		< 0.2	
MRR (nm/min)	73.4		< 9	

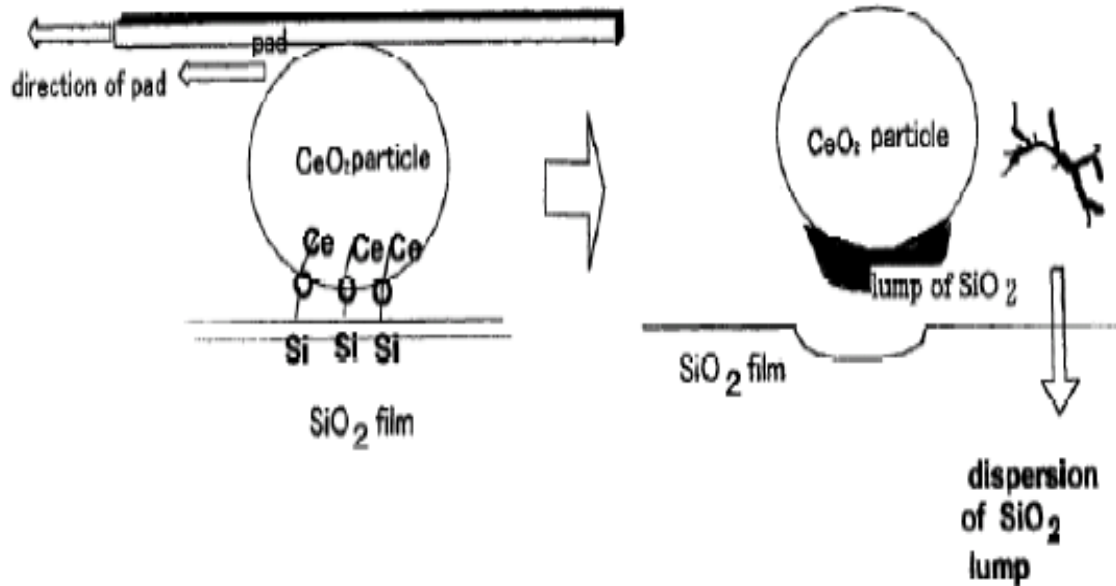
**Colloidal silica also shows a similar behavior**

# Cu CMP results using silica or alumina with 1% glycine and 1% H<sub>2</sub>O<sub>2</sub>

Location found	Fumed Silica		Alumina	
	Cu detected (mg)	% Cu detected	Cu detected (mg)	% Cu detected
Solution	2.2	<b>47.8</b>	16.6	<b>88.3</b>
Abrasive	2	<b>43.5</b>	1.5	8
Pad	0.4	8.7	0.7	3.7
Total	4.6	100	18.8	100
Cu loss from disk per polish	5.1		20.8	
MRR (nm/min)	429		1750	

**Colloidal silica also shows a similar behavior**

# Polishing Mechanisms of SiO<sub>2</sub> with Ceria

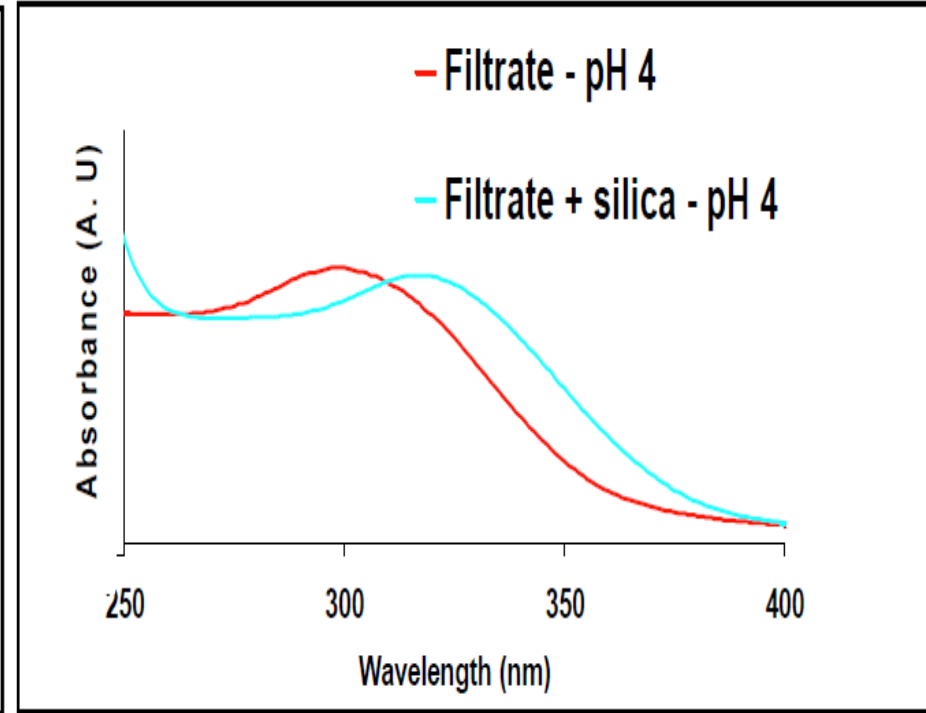
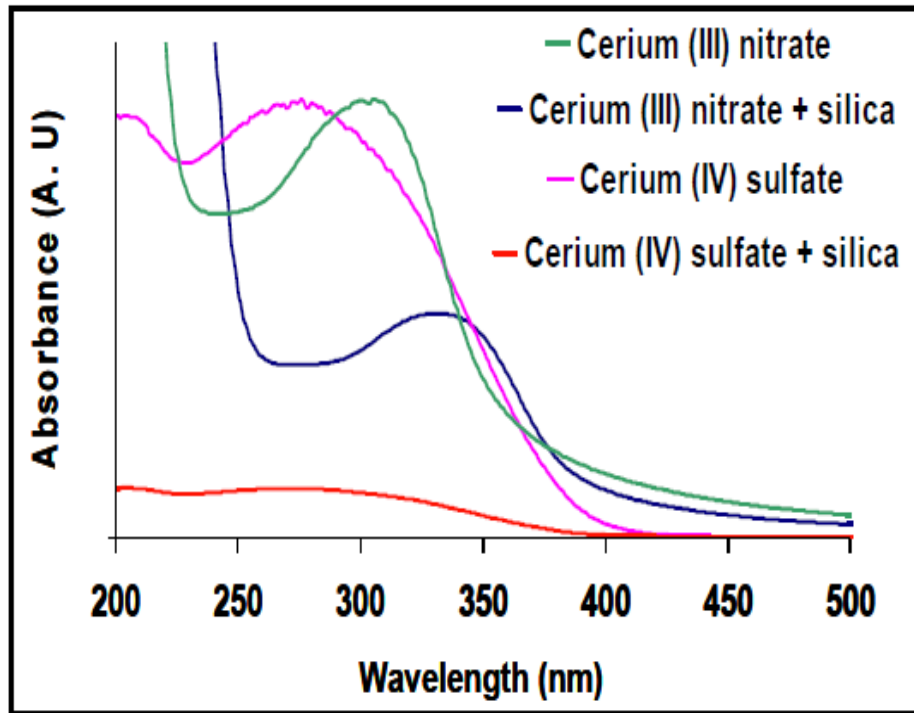


## Using Ceria to Polish SiO<sub>2</sub>

Polishing mechanisms give an idea of the possible species on the surface of nanoparticles in the slurry waste.

- Lee Cook, *Journal of Non-Crystalline Solids* 120 (1990) 152-171
- Tetsuya Hoshino, Yasushi Kurata, Yuuki Terasaki, Kenzo Susa, *Mechanism of polishing of SiO<sub>2</sub> films by CeO<sub>2</sub> particles. Journal of Non-Crystalline Solids* 283 (2001) 129-136

# UV-Vis Absorbance (Reactivity of $\text{Ce}^{3+}$ salt)



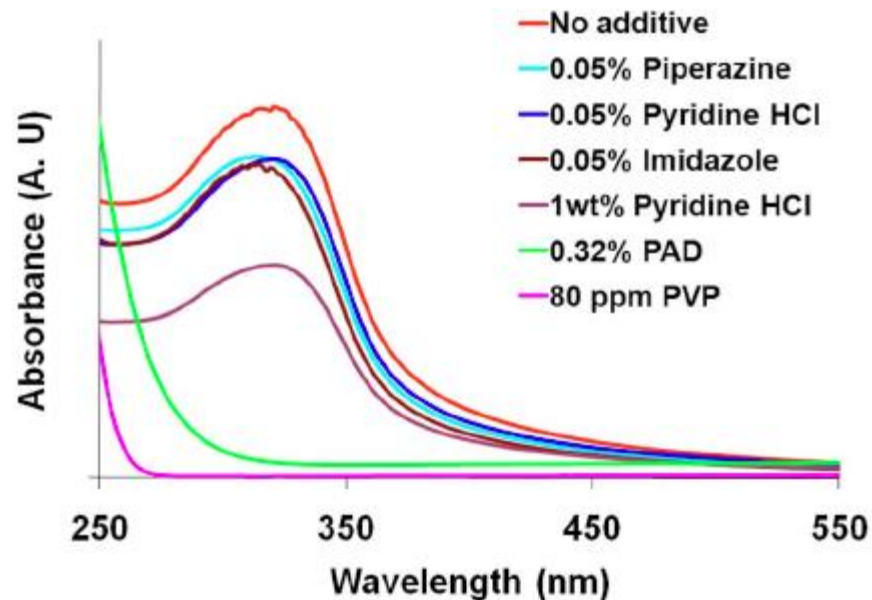
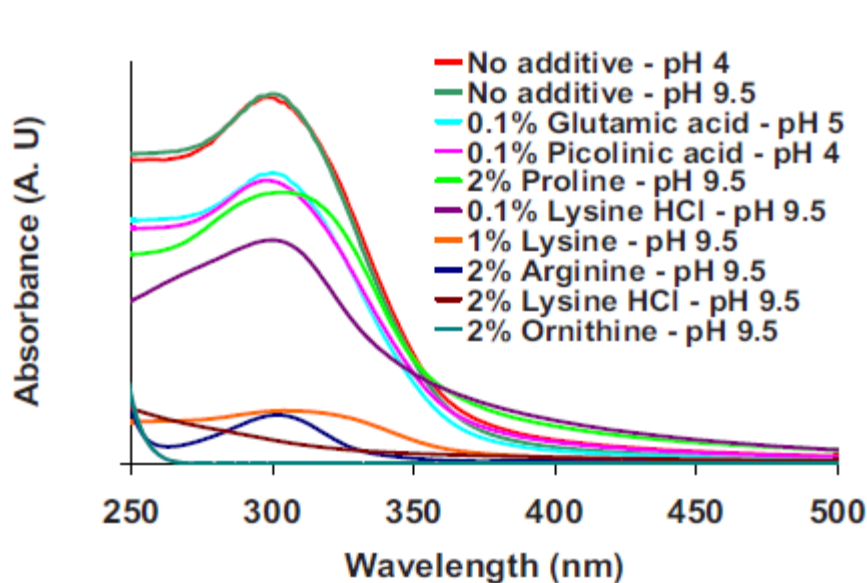
This shift in the peak corresponds to **Ce-O-Si** structures



Ref: Veera Dandu, PhD. thesis. Clarkson University



# Interaction of ceria with different additives

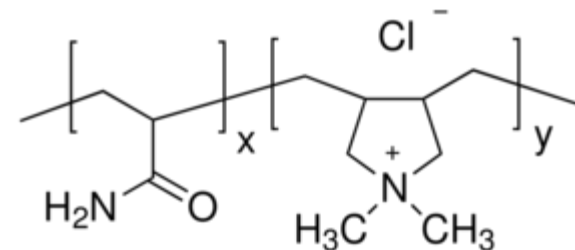
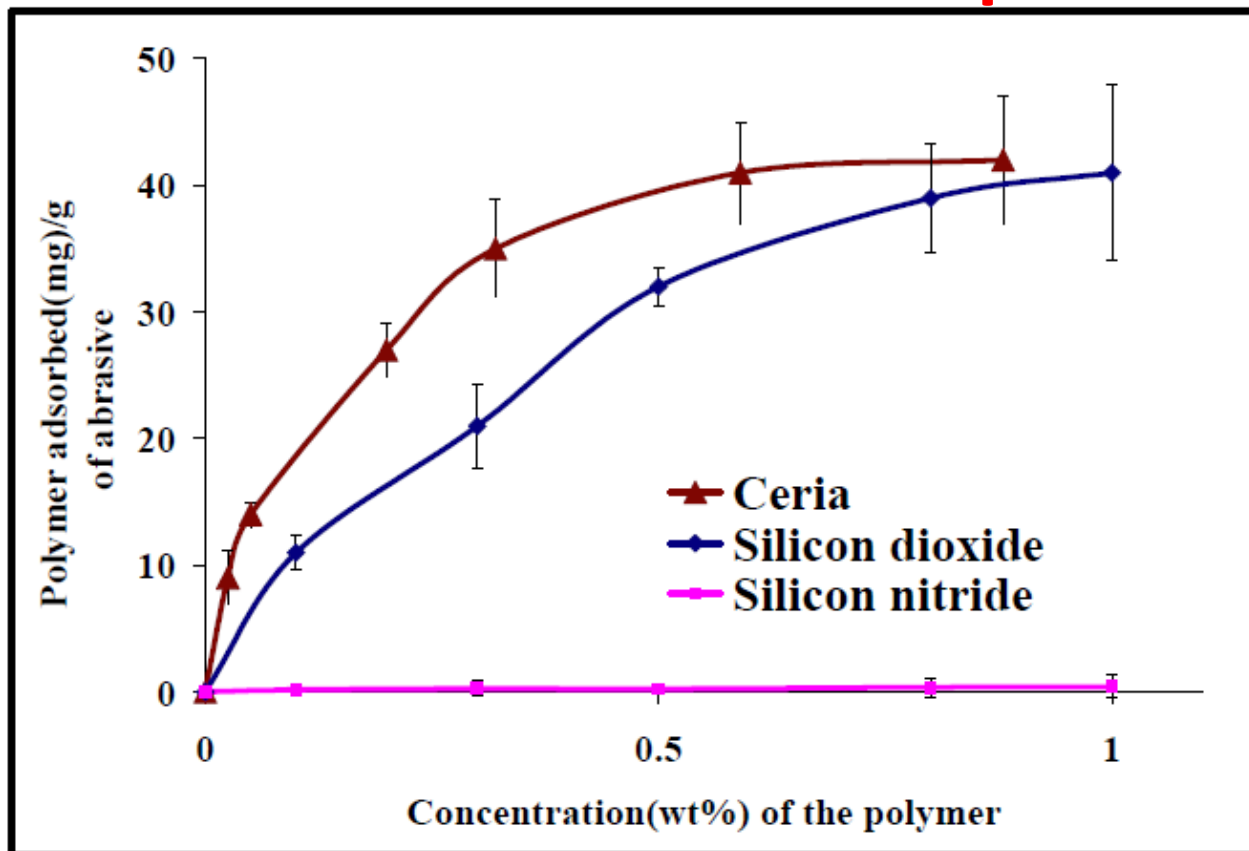


- The RRs of Oxide with the slurries having lower peaks were low (<2 nm/min).
- The absence of the absorbance peak is due to the blocking of all the  $Ce^{+3}$  species on the surface of the ceria particles



Ref: Veera Dandu, PhD. thesis. Clarkson University

# Adsorption Isotherms showing the adsorption of “PAD” on different particles at pH 4



**Poly (acrylic acid-co-diallyl-dimethyl ammonium chloride) (PAD)**

Dispersions : 0.25wt% ceria ( $d_m \sim 60$  nm), 10wt% silicon dioxide ( $d_m \sim 50$  nm) and 10wt% silicon nitride ( $d_m \sim 50$  nm)

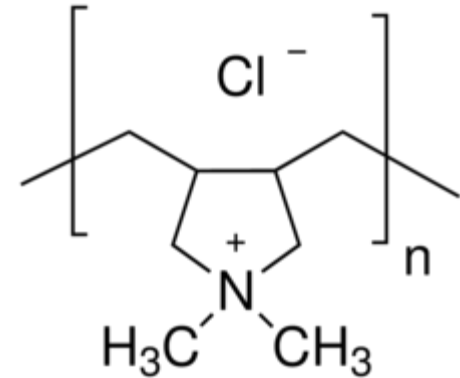
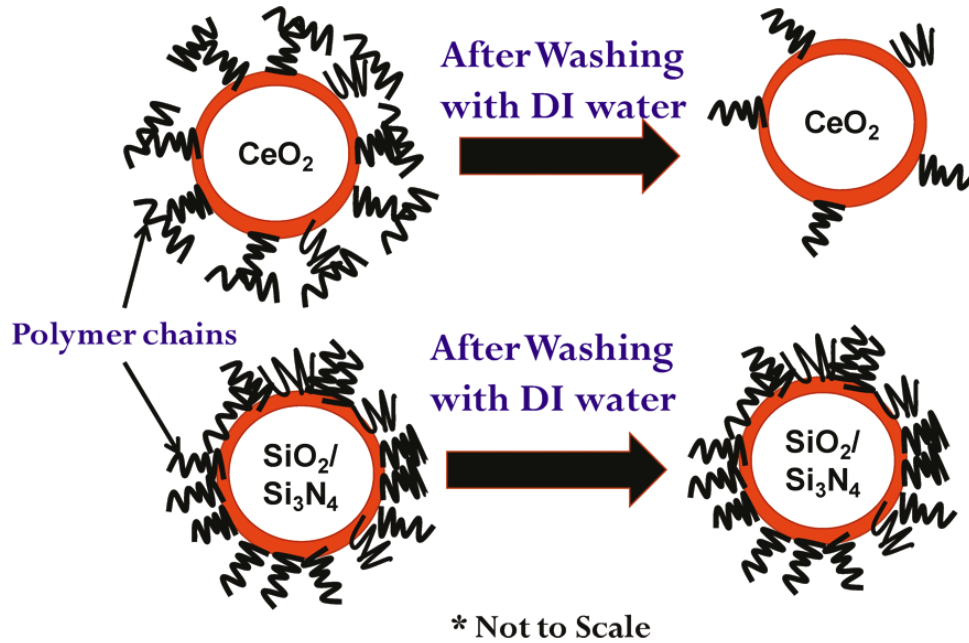


Ref: Veera Dandu, PhD. thesis, Clarkson University

# Effect of washing with water on adsorbed PA and proline at pH=5

No of times redispersed	PA (mg/g of solid) remaining on			Proline (mg/g of solid) remaining on		
	Si <sub>3</sub> N <sub>4</sub>	SiO <sub>2</sub>	CeO <sub>2</sub>	Si <sub>3</sub> N <sub>4</sub>	SiO <sub>2</sub>	CeO <sub>2</sub>
0	96	74	13	147	98	8
1	8	9	4	6	11	1
2	<1	<1	3	1	<1	1

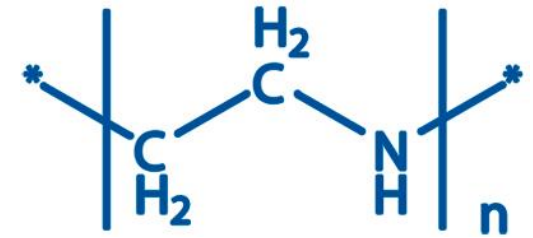
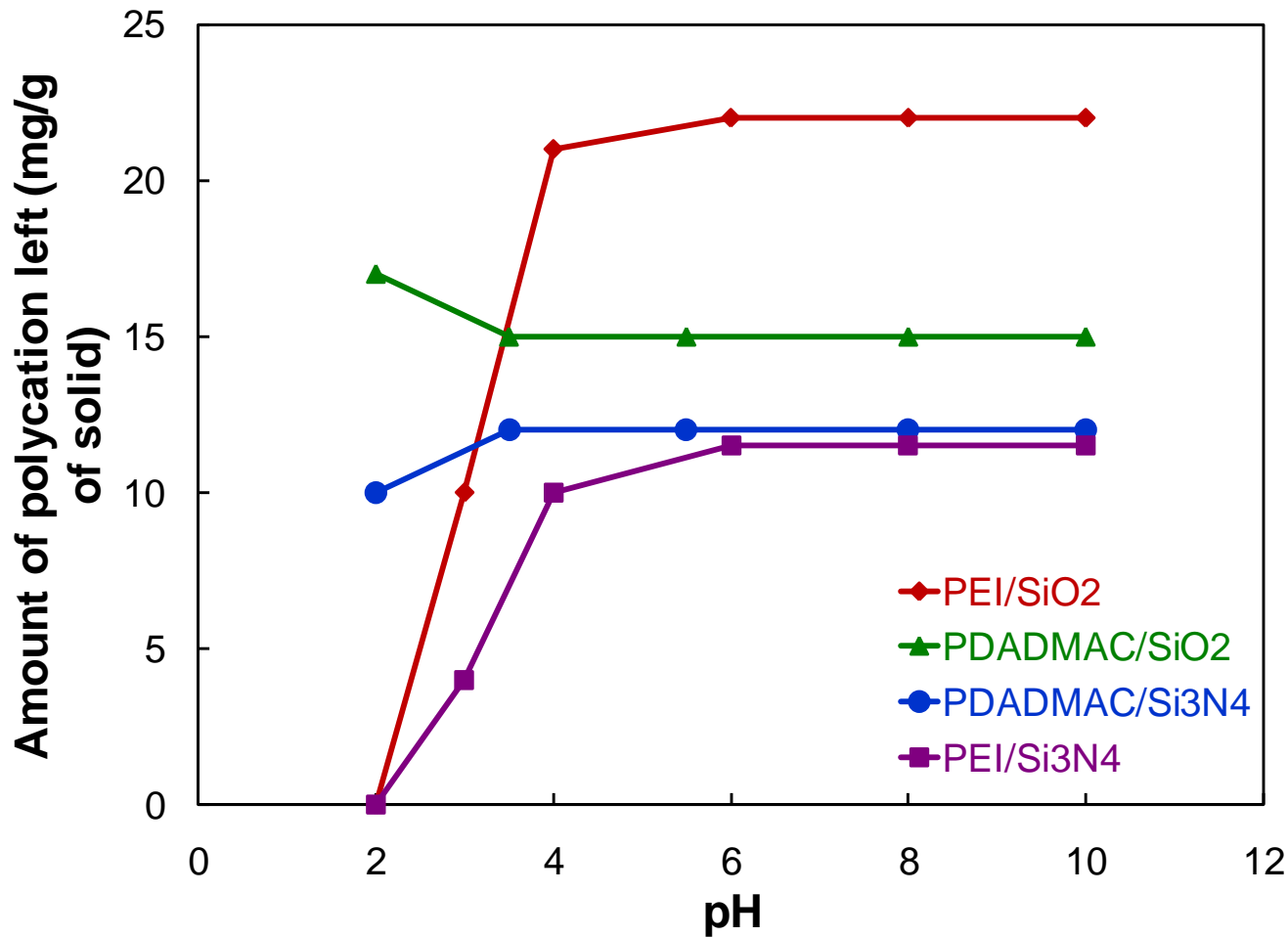
# Adsorption of polymers on particles



Structure of Poly diallyl dimethyl ammonium chloride (PDADMAC)

	Zeta Potential at pH 10		
	Bare	With 250 ppm PDADMAC	Modified particle dispersion
Ceria	-35	+56	+2
Silica	-60	+28	+21
Silicon Nitride	-38	+23	+24

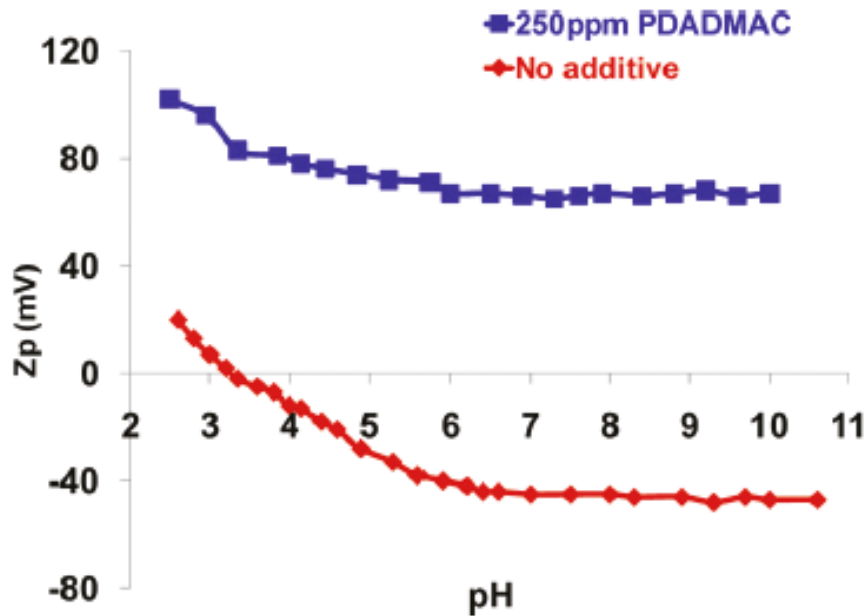
# Adsorption of poly-cations on $\text{SiO}_2$ and $\text{Si}_3\text{N}_4$ surfaces



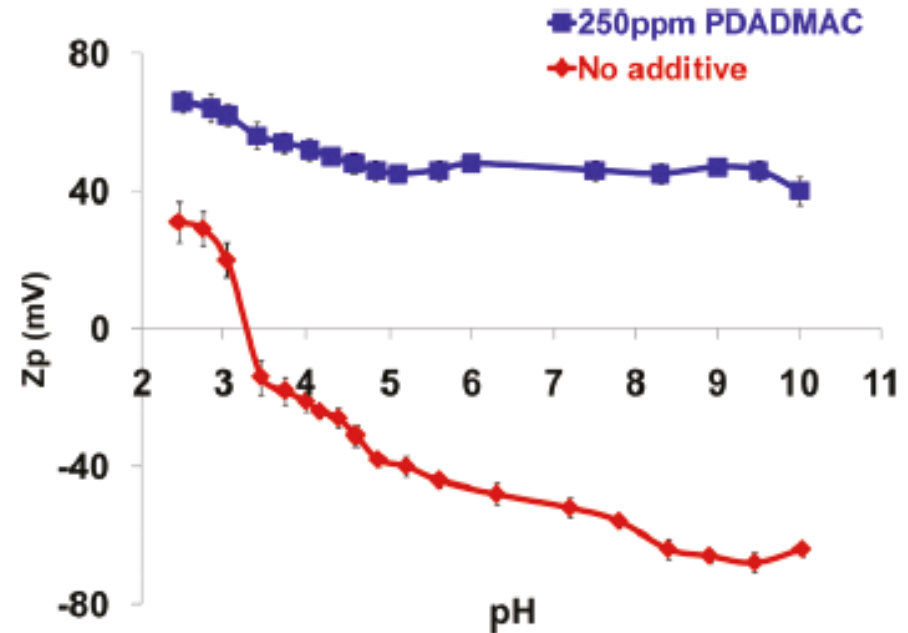
Polyethyleneimine  
(PEI)

Amount of PDADMAC/PEI left on  $\text{SiO}_2$  and  $\text{Si}_3\text{N}_4$  surfaces after 2 washings with DI water at the respective pH values.

# Adsorption of cationic polymers on poly-Si and IC-1000 pad surfaces

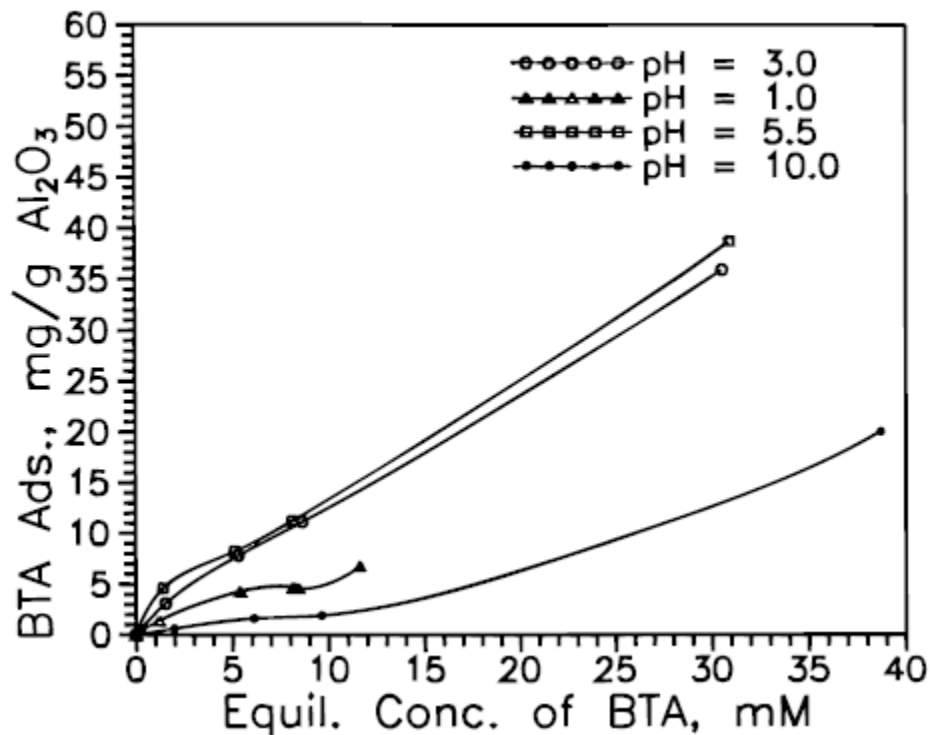


Zeta potentials of IC1000 pad in the absence and presence of 250 ppm PDADMAC.

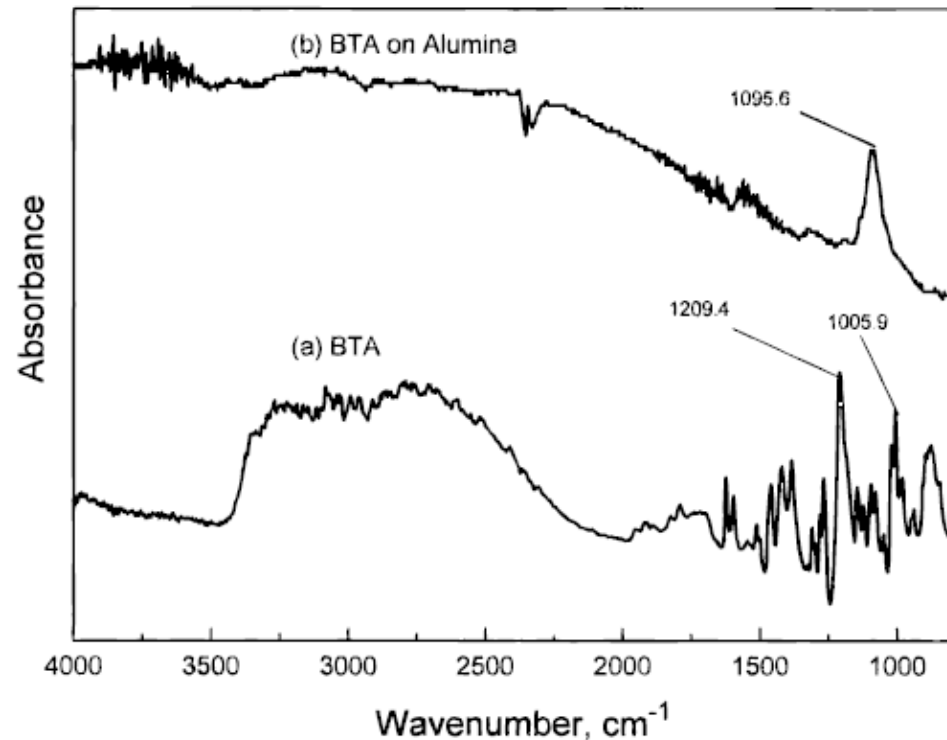


Zeta potentials of poly-Si films in the absence and presence of 250 ppm PDADMAC

# Adsorption of BTA on Alumina



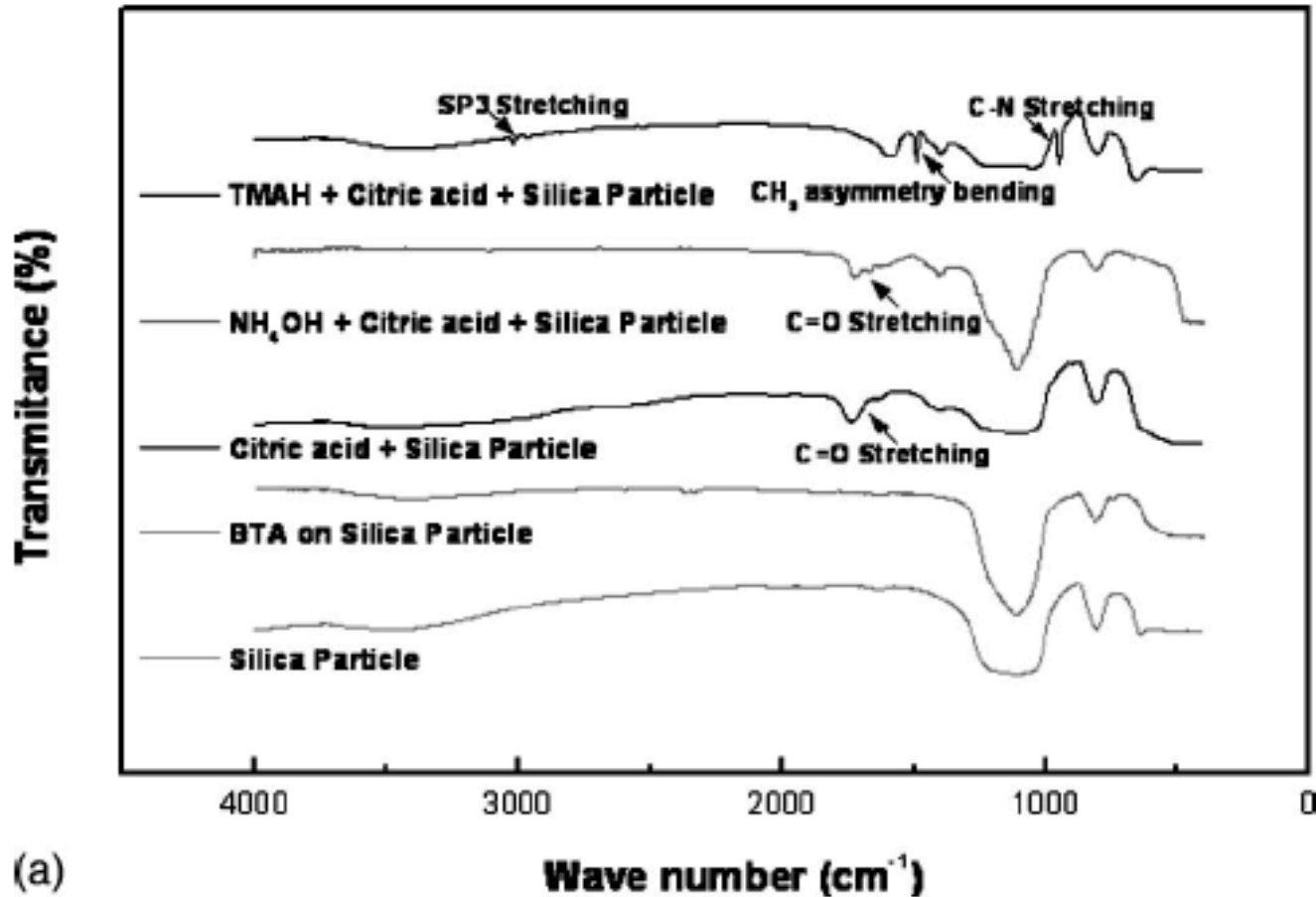
Adsorption isotherms of BTA on alumina particles at various pH's.  
Changes in the adsorption of BTA can modify the surfaces and change the RRs.



## FTIR Spectra:

C-H (stretching) 3053-3090 cm<sup>-1</sup>  
N-N (bending) 1209 cm<sup>-1</sup>  
N-H (stretching) 2800 cm<sup>-1</sup>

# Binding of PCMP additives on silica surfaces

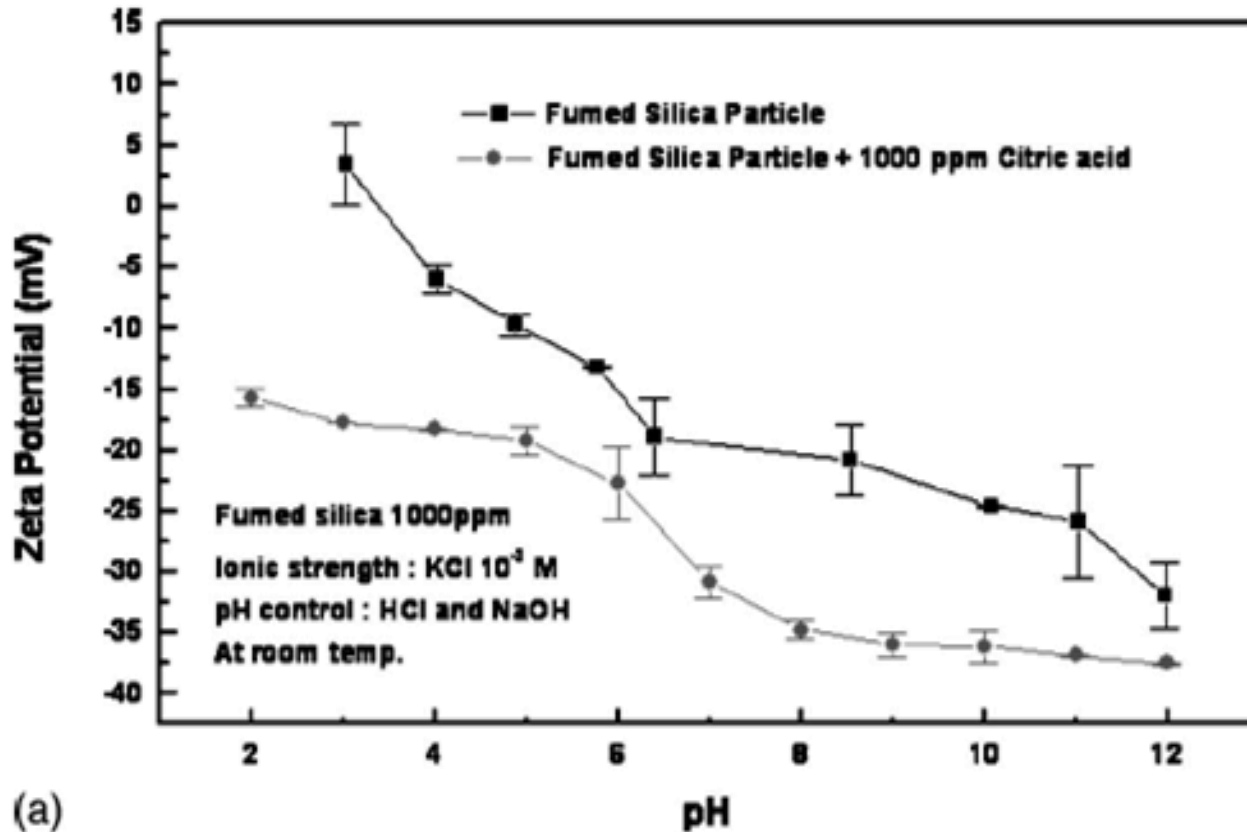


FTIR spectra of silica particles in various cleaning solutions

- Yi-Koan Hong, Dae-Hong Eom, Sang-Ho Lee, Tae-Gon Kim, Jin-Goo Park and Ahmed A. Busnaina. *J. Electrochem. Soc.* 2004, Volume 151, Issue 11, Pages G756-G761



# Zeta Potential of silica surfaces during PCMP cleaning



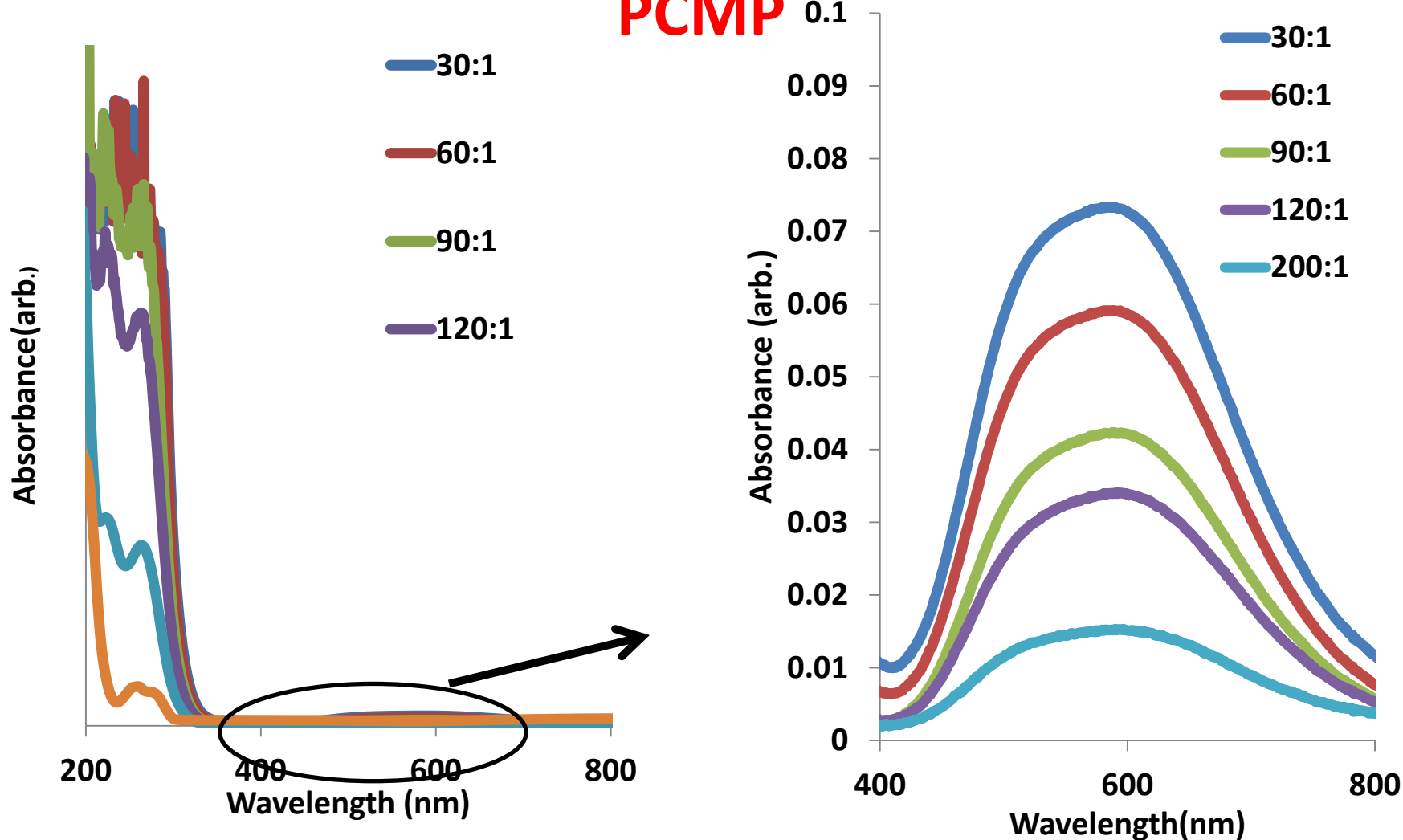
(a)

The zeta potential of (a) silica as a function of pH with and without the addition of citric acid.

- Yi-Koan Hong, Dae-Hong Eom, Sang-Ho Lee, Tae-Gon Kim, Jin-Goo Park and Ahmed A. Busnaina. *J. Electrochem. Soc.* 2004, Volume 151, Issue 11, Pages G756-G761

# UV-Vis of spent 'Planar Clean' solution during Cu

PCMP



**Cu Oxide-Amine complex particles formed due to undercut process during PCMP.**

# Characteristics of GaAs processing waste streams

Source	Characteristics	pH	Total GaAs* (g/L)	Dissolved As (g/L)
Wafer polishing and backside polishing	Clear solution with alumina and SiO <sub>2</sub> particles. May be alkaline or acidic	10-11  6	3-5	1.8-2.4

\* Solid GaAs Particulates and dissolved species (H<sub>3</sub>AsO<sub>3</sub> (pH 2-9)), H<sub>3</sub>AsO<sub>4</sub>)

- <http://www.semiconductor-today.com/features/Semiconductor%20Today%20-%20Management%20of%20arsenic-rich%20waste.pdf>

# Where do Ga and As end up during polish of GaAs?

pH	Element composition ( $\mu\text{g/g}$ )			
	Solids		Liquid	
	Ga	As	Ga	As
2	39.2	33.8	19.8	18.6
10	51.9	47	8.2	26

# Hazards associated with GaAs processing waste streams

Waste Stream	Possible Hazards	Hazard Minimization
Solid particles as dust	<ul style="list-style-type: none"> <li>• Inhalation of dust</li> </ul>	<ul style="list-style-type: none"> <li>• Minimize dust generation through wet processing.</li> <li>• Ventilation and extraction of working area of grinders and saws.</li> <li>• Store waste in liquid slurry form in a closed container.</li> <li>• Wear Protective clothing , mask and gloves</li> </ul>
CMP, grinding, cutting and lapping waste slurries	<ul style="list-style-type: none"> <li>• Ingestion of GaAs.</li> <li>• Dermal Contact.</li> <li>• Contamination of ground water through improper disposal.</li> <li>• Ingestion of Arsenic.</li> <li>• Generation of arsine gas during polish and in waste container.</li> </ul>	<ul style="list-style-type: none"> <li>• Treat waste streams as hazardous waste and dispose of according to local and national regulations.</li> <li>• Wear protective clothing and gloves.</li> <li>• Avoid creating a reducing environment in waste slurry tank.</li> <li>• Maintain low pH in waste slurry to avert microbial activity.</li> </ul>

- <http://www.semiconductor-today.com/features/Semiconductor%20Today%20-%20Management%20of%20arsenic-rich%20waste.pdf>

# Effect of Aqueous Phase pH of Slurries on PH<sub>3</sub> Generation (30 sec polish of InP)

Slurry	Maximum PH <sub>3</sub> concentration (ppb) as a function of pH						
	2	4	6	7	8	10	12
0.3 M aq. H <sub>2</sub> O <sub>2</sub> + 3 wt % silica	140 (34)	135 (4)	94 (10)	73 (16)	n.d.	n.d.	n.d.
0.3 M aq. H <sub>2</sub> O <sub>2</sub> + 3 wt % silica + 0.08 M OA	109 (14)	87 (6)	15 (3)	n.d.	n.d.	n.d.	n.d.
0.3 M aq. H <sub>2</sub> O <sub>2</sub> + 3 wt % silica + 0.08 M TA	120 (12)	69 (8)	26 (5)	11 (2)	n.d.	n.d.	n.d.
0.3 M aq. H <sub>2</sub> O <sub>2</sub> + 3 wt % silica + 0.08 M CA	179 (15)	139 (10)	47 (3)	47 (4)	n.d.	n.d.	n.d.

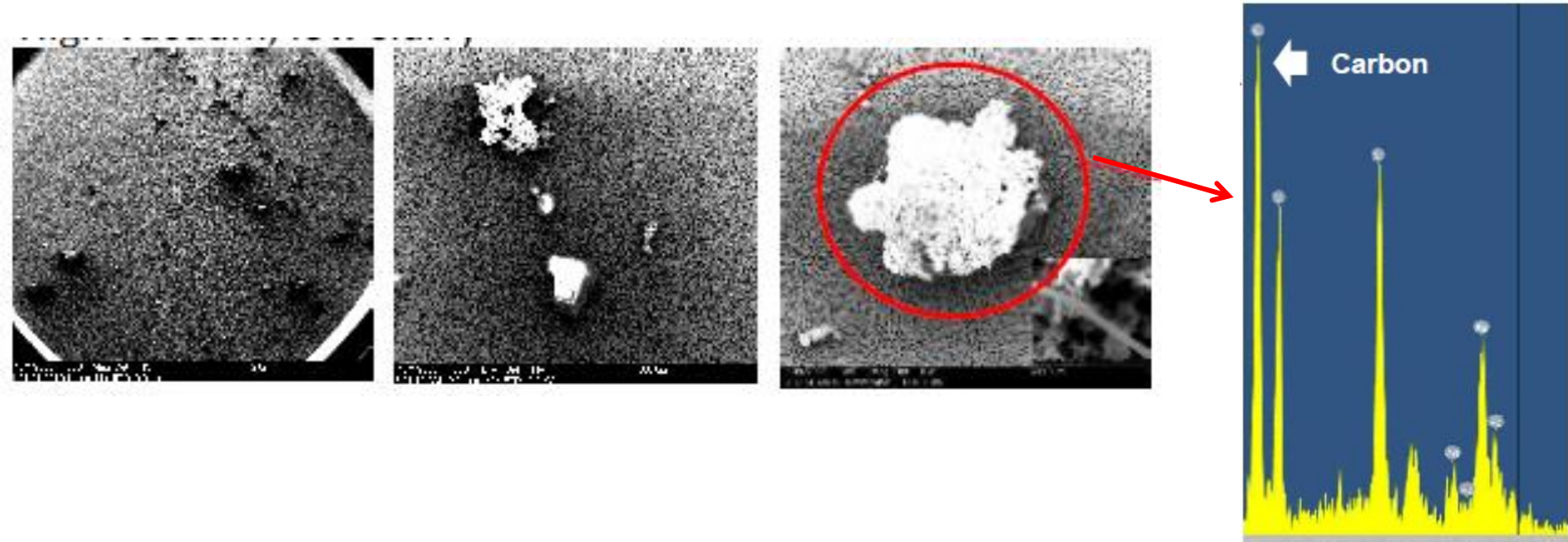
Values in parentheses are standard errors of measurements; n.d. = not detected

**CM-4 gas detector**

# Pad Debris

- Pad debris is often observed during polishing with in-situ conditioning.
- The polyurethane polishing pads possess both –COC- and -COOC - functional groups in the main chain structure that experience some degree of hydrolytic attack with usage.
- Alkaline media accelerates hydrolytic attack and in the presence of weak bases degrades the polyurethane more rapidly than neutral water.

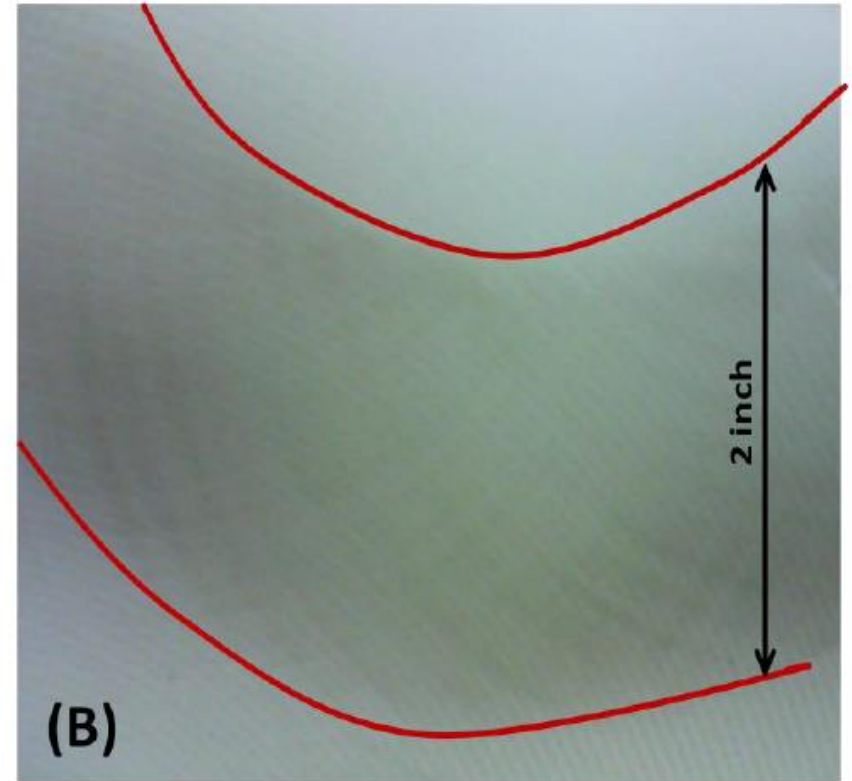
# Pad Waste generated during CMP



- Particles in the effluent included slurry, agglomerates, and pad debris
- Carbon peak from SEM EDS identified pad debris

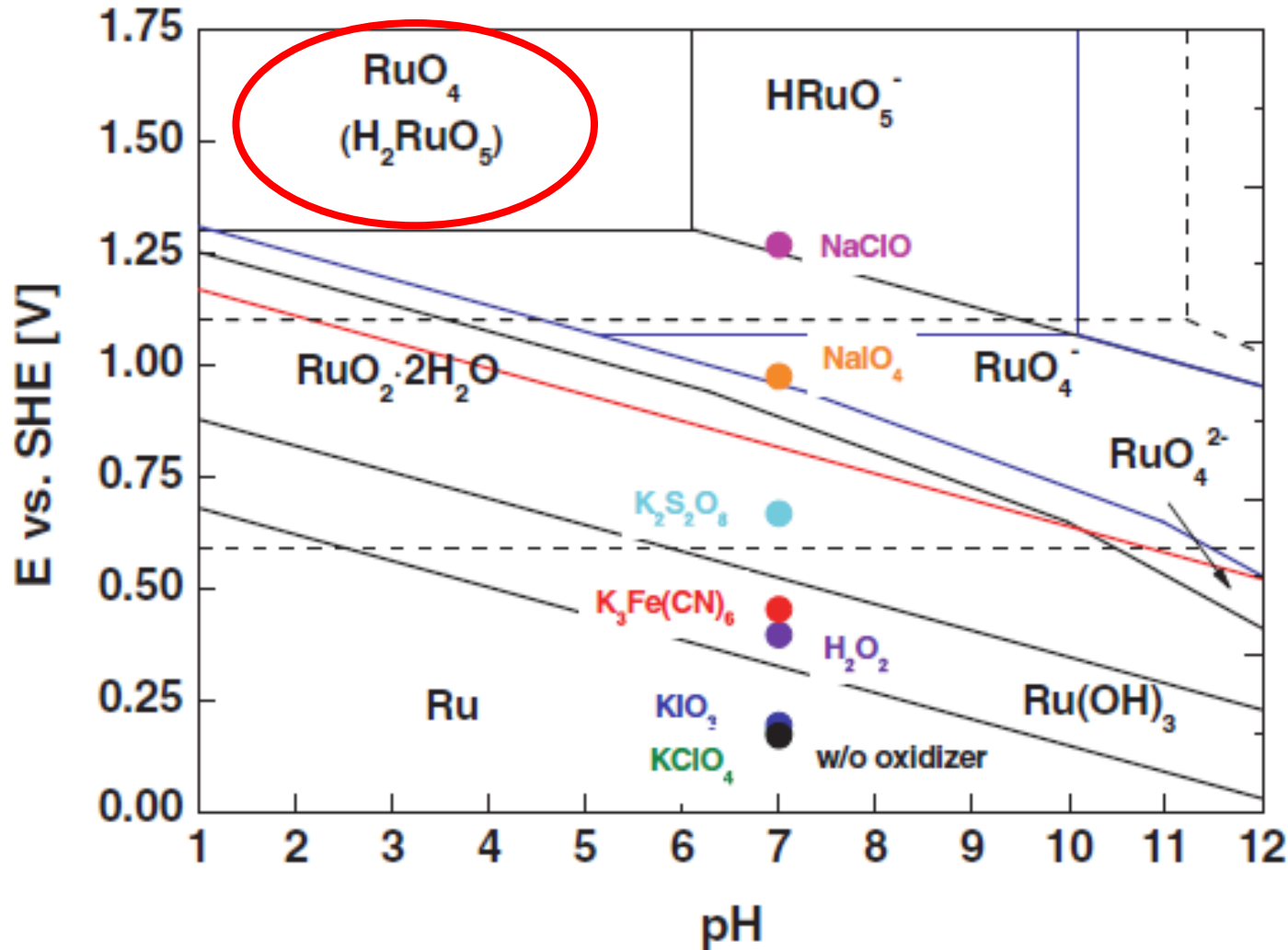


# Pad staining during Ru polishing with $\text{KIO}_4$



Polishing pad (IC 1000) before (A) and after (B) polishing with a slurry containing 0.015 M  $\text{KIO}_4$  + 5 wt% Silica at pH 6

# Issues with Ruthenium CMP

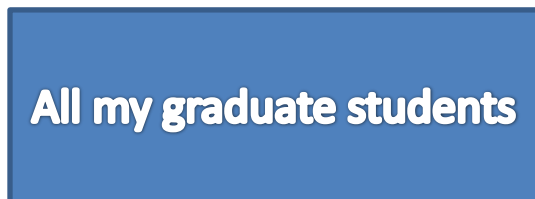
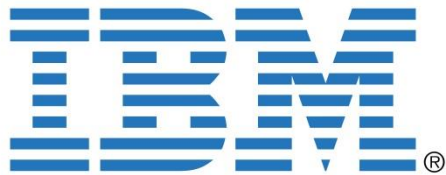


$\text{RuO}_4$  :

Volatile, toxic and explosive

- Hao Cui, Jin-Hyung Park and Jea-Gun Park *ECS Journal of Solid State Science and Technology*, 2 (1) P26-P30 (2013).

# Acknowledgments



Thank You