



AES NEWSLETTER



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Many thanks to our supporters and friends for their generous contributions.

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GE Healthcare

LabSmith

Kendrick Labs, Inc

Our traditionally strong meetings would simply not be possible without help from our supporters. Their donations are greatly appreciated.

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POWER UP AES MEMBERS, 2011 HAS BEGUN!

Welcome Message from the New AES President

As we start 2011, the American Electrophoresis Society is as strong as ever and growing rapidly! Research in linear and nonlinear electrophoretic applications especially in the fields of microchips and nanotechnology have exploded in the last decade. The overlap of electrokinetic tools and complex biological systems has opened up whole new analysis capabilities. What an exciting time to be an Electrophores-er!

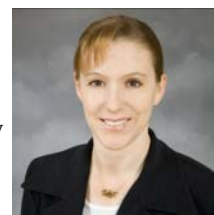
The council has begun efforts to expand our strong programming into AES special symposia at related meetings in order to better serve our diverse membership. We hope in the next two years to reach out to: the microfluidics community via Chemical and Biological Microsystems Society (CBMS)'s annual mTAS conference (<http://www.microtas2011.org>), the chemical and spectroscopic community via the Federation of Applied Spectroscopy Societies (FACSS) annual conference (<http://facss.org/facss/index.php>) and the biological engineering community via Biomedical Engineering Society (BMES, <http://www.bmes.org>).

In addition, the AES council has refocused committees to a) launch a membership drive, b) begin an annual honorary award with corresponding symposia in honor of AES pillars as well as current research superstars, and c) expand our sponsorship. We are ever grateful to our loyal annual sponsors such as Bio-Rad Laboratories, CBS Scientific, Decodon, Kendrick Labs and GE Healthcare as well as new sponsors LabSmith and COMSOL. By fostering and growing strong interactions with industry, AES will facilitate electrophoresis related research and applications.

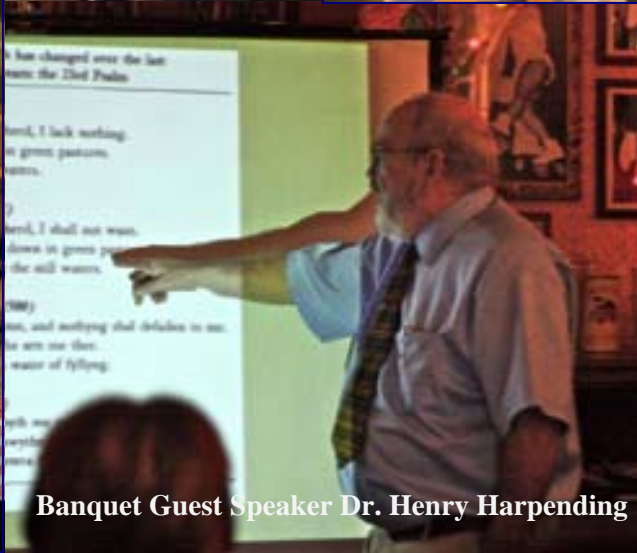
AES is also in the process of partnering with the journal *Electrophoresis* to do a special issue for our 2011 annual meeting held jointly with AIChE. We are starting a student chapter so please share this newsletter with your students and have them look us up on Facebook and LinkedIn.

As always, if there is anything the council or I can do to help facilitate your work in this "electrifying" field, please do not hesitate to contact us.

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2010 Meeting Moments:



Banquet Guest Speaker Dr. Henry Harpending



Winners of the 2010 Poster Competition: (right to left)

First Place: Sachidevi Puttawsamy
Ipod
University of Missouri
Biological Engineering

Second Place: Alice Jernigan
(\$75)
University of Arkansas
Chemical Engineering

Third Place: Paul Jones
(\$25)
Arizona State University
Chemical Engineering



Determining Dielectrophoretic Mobility in DC Insulator-based Systems by Mark A. Hayes, Noah S. Weiss, Prasun Mahanti, and Paul Jones, Chemistry and Biochemistry Department, Arizona State University

Particles are ubiquitous in our bodies and our environment. This class of materials includes cells, organelles, nanoparticles, aerosols, large proteins and DNA strands, bacteria, and viruses—among other organic and inorganic debris. As the members of AES know, dielectrophoresis (DEP), specifically insulator-based DEP (iDEP), is continuing to emerge as an important technique for manipulating these micro- to nano-scale particles. Within these important works, comparison of theoretical constructs to real-world data is only beginning. One limitation in connecting the properties of particles with dielectrophoretic (and electrokinetic) actions is a lack of valid physical descriptions based on first principles. Currently, assessments are made that ignore any subtle particle properties, or limit the discussion to polarizability and/or the Clausius-Mossotti factor. In fact, most discussions of real data rarely advance beyond an analysis of positive (towards stronger fields) versus negative (towards weaker fields) DEP.

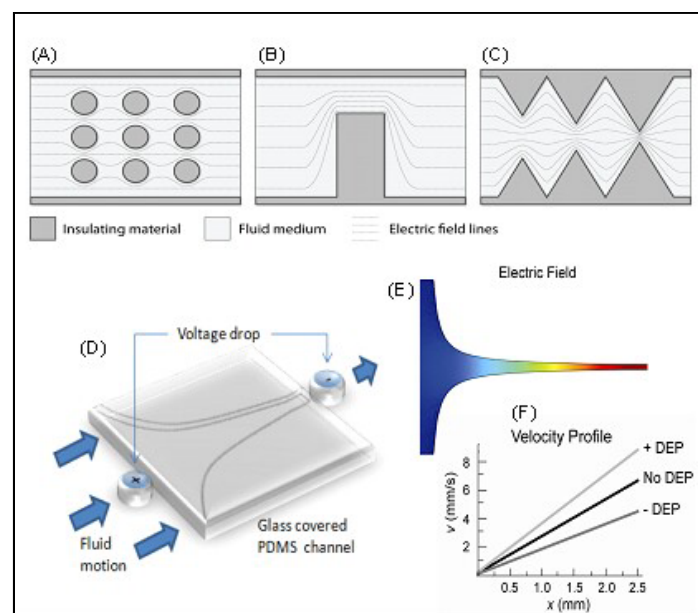
Another approach is to define a DC-iDEP mobility according to the form of electroosmotic and electrophoretic mobilities. The DEP mobility (μ_{DEP}) is defined independent of the electric field gradient and thus becomes a universal parameter. In other words this definition of DEP mobility is intrinsic to the particle and represents the relative DEP velocity per unit electric field gradient squared as shown: and, where v_{DEP} is the DEP velocity, E is the electric field, ϵ_f is the permittivity, r_p is the particle radius, $Re(fcm)$ is the real part of the Clausius-Mossotti factor defined by the particle and medium conductivities (σ) at low frequency (f) and η is the fluid viscosity. Furthermore, an ideal method for quantifying iDEP ought to simultaneously quantify other electrokinetic effects (electroosmosis and electrophoresis).

We have initiated a strategy to quantitatively determine dielectrophoretic mobility in a converging microfluidic channel where dielectrophoretic and electrokinetic mobilities of polystyrene particles are determined simultaneously using a unique streak-based velocimetry to generate the velocity profile of particles (see figure). The electrokinetic mobilities are determined in the non-converging zone and the funnel-shaped section has a constant gradient—allowing for trivial differentiation of DC-iDEP actions. An automated algorithm detects, processes, and determines the velocities traveled by the particles at all pixels across the image sequence. Streak-based velocimetry operates by associating the length of a particle trajectory (streak) with the exposure time to estimate the velocity field.

Using 1 μm polystyrene particles the electrokinetic mobility was estimated to be $3.5 \times 10^{-4} \text{ cm}^2/(\text{V}\cdot\text{s})$ using the velocity data from non-converging zone and the dielectrophoretic mobility was $-2 \times 10^{-8} \pm 0.4 \times 10^{-8} \text{ cm}^4/(\text{V}^2\cdot\text{s})$ ($n=3$) was determined from the converging zone using the slope of the velocity profile. This result agrees with the general finding that polymeric particles exhibit negative dielectrophoresis under similar conditions. Under these particular conditions the electrokinetic velocity is

about twice the magnitude of the dielectrophoretic velocity. The relative DEP velocity could be increased by increasing the applied voltage, reducing electroosmotic flow, or utilizing steeper field gradients. However, particle motion rapidly changes direction in these regimes complicating velocity associations. For optimal quantitative analysis, the DEP motion must be observable but not predominant.

Quantitative approaches like this one enable an unprecedented evaluation of iDEP and provide a metric for standardization. Ideally, discussions will eventually evolve from subjective descriptions of particle behaviors to more objective quantitative responses.



Schematic showing A)-C) various forms of iDEP patterns, D) 3-D representation of the converging channel design, E) calculated electric field in channel, and F) velocity profiles for no, positive and negative DEP forces with the channel. Design allows for determination of particle properties for use as probes for A)-C) systems.



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We are pleased to announce that the 2011 AES Meeting will be held on October 16-21 at Minneapolis, MN as an AIChE Topical Conference. **Mark your calendars!**

The 2011 AES meeting will be held at the Minneapolis Convention Center. With nearly 480,000 square feet of trade show space, 87 column-free conference meeting rooms, a 28,000-square-foot ballroom, and an auditorium with superb production and flexible technology options, the facility can handle any event from a small meeting to a large convention. Conveniently located close to freeways and only 20 minutes from the airport on the southern edge of downtown Minneapolis, the Convention Center is easy for attendees and exhibitors to find. Plus, the convention center is connected by climate-controlled skyways to the downtown hotels, restaurants and shopping. For more information please visit www.minneapolisconventioncenter.com



#	Session Description
1	Advances In Electrokinetics and Electrophoresis - Fundamentals
2	Microfluidics: Bioanalytical Applications
3	Plenary Session of the American Electrophoresis Society
4	Microfluidics-- Detection
5	Fundamentals of Electrokinetic Flows: Novel Applications and Ionic Fluxes at Interfaces
6	Biomedical Diagnostics I
7	Nanoscale Electrokinetics
8	Electrokinetic Behavior of Microparticles and Nanoparticles: Fundamentals and Applications
9	DNA Analysis in Microfluidic and Nanofluidic Devices
10	Poster Session for the American Electrophoresis Society
11	Advances in Electrophoretic Protein Separation and Analysis
12	Electroporation and Electrophysiology
13	Award Session of the American Electrophoresis Society

The call for papers (PTP, Proposal To Present) for the 2011 AES meeting is now open! **We are excited to announce the addition of 4 new microfluidic and protein separation sessions this year!** The program is sketched out in the table above. We are still looking for a few session co-chairs so please be sure to contact us if you're interested. Questions are welcome.

Please note:

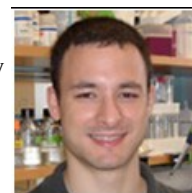
The Call for Papers

- ◆ Opens – Jan. 17, 2011
- ◆ Closes – May 2, 2011

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2011 Meeting Co-Organizers

**Election Results:
Our Three New
Councilors,
pictured to the
right, were
welcomed at the
2010 board meet-
ing in Salt Lake
City.**



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