

Topics

- Electrochemical Detection
 - Capillary electrophoresis.
 - Electrochemical impedance spectroscopy (EIS) See Gamry Instrument article
- Gamry Instrument article

 Optical Detection

 Conventional Off-Chip or "Free-Space"

 Absorbance Detection
 Fluorescence Detection
 Chemiluminescence
 Surface Plasmon Resonance
 Surface Enhanced Raman
 Spectroscopy

- On-Chip Methods
 - Frequency Specific LEDs
 - Absorbance Spectroscopy
 - Fluorescence
 - Chemiluminescence
 - Optical Waveguides
 - Surface Plasmon Resonance
 - Interferometry
 - Holography

Glass and Polymer Based Devices













Positive "Downscaling" Effects

- Reduced sample and reagent volumes.
- Ten fold reduction in length scale.
- Laminar flow (low Reynolds numbers).
- Improved mass flow by diffusion.
- Harvesting electrokinetic effects
- The surface-to-volume ratio (SVR) may increase by a factor of more than ten thousand during downscaling and solute/wall interactions become dominant.

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a) Sample introduction and electrophoretic separation are accomplished in each of two crossing channels. b) Sample is driven through the short sample channel across the separation channel by application of a potential, and c) The introduced sample "plug" (nanoliter volume) is then electrophoretically separated by application of another potential.

A. High voltage along the length of the capillary separates the analytes and drives them toward the detection electrodes. B. A constant potential is applied with a potentiostat to the detection electrodes. C. As each species passes, a change in current or conductivity between the detection electrodes is detected. Stevent 8. Salterman Concogue, A and Adresser R. Morents. Electrodemical detection lechniques in micro- and narofluidic devices.

Video of Capillary Electrophoresis... Capillary Electrophoresis (CE) https://youtu.be/wStV1rFjHOo

Recall Electrochemical Impedance Spectroscopy (EIS)...



(For a more detailed discussion of EIS see the Gamry Industries

- a) Three-electrode electrochemical cell. (WE = Working Electrode, CE = Counter Electrode, RE = Reference Electrode & V = Voltage).
- b) On the left, a small AC voltage is applied across the WE and RE of a three-electrode electrochemical cell.
 c) Current is then measured from the WE to the CE, and varies with frequency
- and the analyte concentration.

Entifation.

3) Gerooglu. A and Adrienne R. Miserick. Electrochemical detection techniques in micro- and nanofluidic devices. Microdict Handball (2014) 17.81-807.

b) Fast I., Zhao G, Shi H, Lis M, Li Z (2013) A highly selective electrochemical impedance spectroscopy-based applications or sensitive detection of acetamipris. Biosens: Bioelectron 43:12-18. doi:10.1016/j.bios.2012.11.033 applications for sensitive detection of acetamipris. Biosens: Bioelectron 43:12-18. doi:10.1016/j.bios.2012.11.033

What is Impedance?

- Impedance (Z) Total opposition a device or circuit offers to flow of an alternating current at a given frequency.
 - Units are in ohms
 - Complex quantity graphically shown on a vector plane (real and imaginary parts): R +

jX where R is resistance and X reactance

Recall Aptamers...

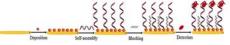
- Aptamers are artificial nucleic acid ligands that can be generated against amino acids, drugs, proteins and other molecules.
- Function similar to antibodies.
- Applications:
 - Therapeutics,
 - Target validation,
 - Drug screening,
 - Affinity separation,
 - Diagnostics and biosensors.



Image courtesy of Archemis

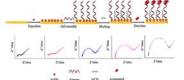
Example of an EIS Implementation...

- Typically applications depend on a specific binding event between the analyte and a recognition element (e.g., enzyme, DNA probe, and antigen), which form a surface layer on the detection electrode, bound directly or through a linker molecule, as shown.
- Shown is modification of an Au working electrode surface for selective detection of the analyte acetamiprid via specific analyte—aptamer binding.



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 The bare Au surface plot is shown leftmost. The spectra changes as the surface is functionalized with Au nanoparticles, aptamer, and a blocking agent to block nonspecific binding sites on the aptamer molecule.



Nyquist plots (impedance in ohm). The extent of the shift in EIS response (two final spectra) upon analyte binding, is a function of the analyte concentration. [See also Bode plots.]

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a) Gencoglu, A and Adrienne R. Minerick. Electrochemical detection techniques in micro- and nanofluidic devic Microfluid Nanofluid (2014) 17:781–807
5) Fan L. Zhao G, Shilt H, Lui, M. Li Z (2013) A highly selective electrochemical impedance spectroscopy-based abalasensor for sensitive detection of acetaminorid. Biosens Bioelectron 43:12–18. doi:10.1016/i.bios.2012.11.03

Optical Detection: Off-Chip

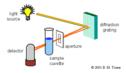
- Absorbance Detection
- Fluorescence Detection
- Chemiluminescence
- Surface Plasmon Resonance
- Localized Surface Plasmon Resonance (LSPR)
 Spectroscopy (electromagnetic-field enhancement that leads to surface enhanced Raman Scattering and other surface-enhanced processes)

teven S. Saliterma

Gai, Hongwei, Yongjun Li, and Edward S. Yeung. 2011. Optical Detection Systems on Microfluidic Chips. Microfluidic Technologies and Applications 304, 171-201.

Absorbance Detection

- UV/Vis light is used for absorption spectroscopy.
- Absorption spectra are related to the structure and concentration of the analyte and are based on the capability of samples to attenuate the incident electromagnetic radiation at various wavelengths.

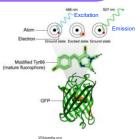


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Introducing Fluorophores...

- A fluorophore is a fluorescent chemical compound that may re-emit light upon light excitation.
- Fluorophores typically contain several combined aromatic groups, or plane or cyclic molecules with several π bonds.
- Generally covalently bonded to a macromolecule, serving as a marker (or dye, or tag, or reporter) for affine or bioactive reagents (antibodies, peptides, nucleic acids).
- Fluorophores are notably used to stain tissues, cells, or materials in a variety of analytical methods.

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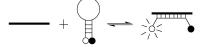


Fluorophore Characteristics

- Maximum excitation and emission wavelength (nm):
 - Corresponds to the peak in the excitation and emission
- Extinction Coefficient (Mol⁻¹cm⁻¹):
 - Links the quantity of absorbed light, at a given wavelength, to the concentration of fluorophore in solution.
- Quantum yield (emitted/absorbed photons):
 Efficiency of the energy transferred from incident light to emitted fluorescence
- Lifetime (in picoseconds):
 Duration of the excited state of a fluorophore before returning to its ground state
- Stokes shift:
 - Difference between the max excitation and max emission wavelengths.

Introducing Molecular Beacons...

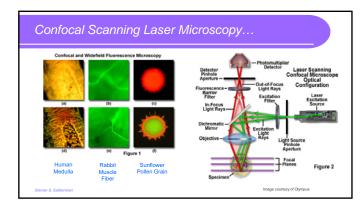
• Single-stranded oligonucleotide hybridization probes that form a stem-and-loop structure:



Target

Molecular Beacon Fluorophore Quencher

Molecular Beacons... Target DNA Fluorophore Molecular Beacon



Chemiluminescence

- Chemiluminescence (CL) is the generation of light (visible, ultraviolet and infrared) by the release of energy from a chemical reaction.
- Advantages:
 - No excitation source (as does fluorescence and phosphorescence),
 - Only a single light detector such as a photomultiplier tube,
 - No monochromator and often not even a filter.
- Detection limits can be 10 to 100 times lower than other luminescence techniques.
- Peroxyoxalates are esters formed by the reaction of hydrogen peroxide and oxalate esters.

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Chasteen, TO

Peroxyoxalate Chemiluminescence

 Peroxyoxalate "light-stick" reaction and emission from the reaction of luminol with hydrogen peroxide in basic aqueous solution catalyzed by cobalt II:





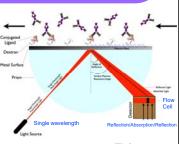


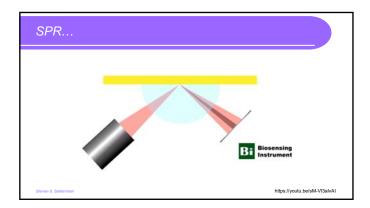
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Image courtesy of Simon W. Lewis, Deakin Universi

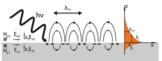
Surface Plasmon Resonance Resonant oscillation of conduction electrons at the interface between a negative and positive permittivity material stimulated by incident light.

Detection is possible because adsorbing molecules cause changes in the local index of refraction, changing the resonance conditions of the surface plasmon waves.





What are Surface Plasmons?



- Coherent delocalized electron oscillations that exist at the interface between any two materials.
- Surface plasmon polaritons are surface electromagnetic waves that propagate in a direction parallel to the metal/dielectric (or metal/vacuum) interface.
 Since the wave is on the boundary of the metal and the external medium (air or water for example), these oscillations are very sensitive to any change of this boundary, such as the adsorption of molecules to the metal surface

Localized SPR Explained...

- Localized surface plasmon resonance (LSPR) spectroscopy of metallic nanoparticles is a powerful technique for chemical and biological sensing experiments
- Materials that possess a negative real and small positive imaginary dielectric constant are capable of supporting a surface plasmon resonance (SPR).
- Plasmonics is the study of these particular light-matter interactions

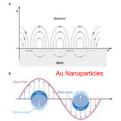
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Willets, Katherine A. And RP Van Duyne . Localized surface plasmon risonance spectroscopy and sensing Annu. Rev. Phys. Chem. 2007. 58:267–97

"Localized" Surface Plasmon Resonance

- Basis for measuring adsorption of material onto planar metal (typically gold and silver) surfaces or onto the surface of metal nanoparticles.
 - Surface plasmon polariton (or propagating plasmon).
 - plasmon).

 Localized surface plasmon. Light interacts with particles much smaller than the incident wavelength.
 - This leads to a plasmon that oscillates locally around the nanoparticle with a frequency known as the LSPR. Similar to the SPR, the LSPR is sensitive to changes in the local dielectric environment.



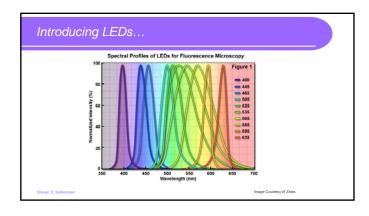
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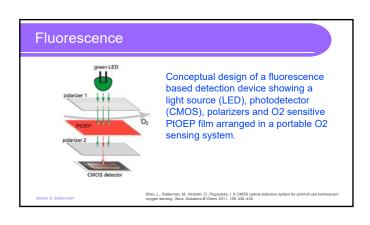
Optical Detection: On-Chip

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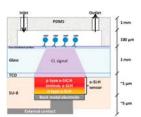
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Absorbance Spectroscopy On-Chip LED 459 minors LED Glass Increasing optical path length by making the light pass axially between parallel 45 degree mirrors. Nota T, Takar H, Vontrola K, Oku N, Ashiki M, Sareada K, Mataumoto K, Ishida M(2006) Serra Actuators B 19245-290 Gai, Horgani F, vongin Li, and Glawed S, Yanag, 2011. Optical Defection Systems on Microfluidic Chips. Microfluidica: Tourhooliges and Applications 304, 117-201.



Chemiluminescence



Integrated opto-microfluidic sensor with a hydrogenated amorphous silicon (a-Si:H) photodetector prepared onto a glass substrate covered by a transparent conductive oxide (TCO) film.

Caputo, D.; de Cesare, G.; Dolci, L.S.; Mirasoli, M.; Nascetti, A.; Roda, A.; Scipinotti, R. Microfluidic chip with integrated a-St:H photodiodes for chemiluminescence-based bioassays. IEEE Sens. J. 2013, 13, 2595–2602.

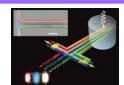
Waveguides for LOC based on Refractive Index

- 1. Solid-state waveguides.
- Solid fibers (silica, glass or polymer) enter the chip and intersect the fluidic channel, exciting analytes and collecting the response.
 Liquid-core waveguides (LCWs).
- The microchannel not only transports the sample, but also transmits the light.
- If liquid (particularly water in a microfluidic chip, with refractive index of 1.33) is used as the core of waveguide, the refractive indices of the cladding should be smaller.
- Liquid-core/liquid-cladding (L²) waveguides

 Two or more different laminar liquids of different refractive index flowing inside a fluidic channel.
- The index of the cladding liquid is smaller than that of the core liquid so that the light is guided in the channel by the total internal reflection.

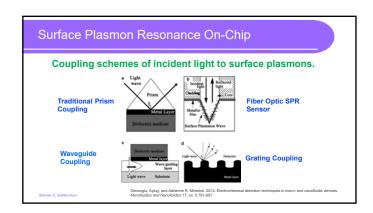
Gai, Hongwei, Yongjun Li, and Edward S. Yeung. 2011. Optical Del Technologies and Applications 304, 171-201.

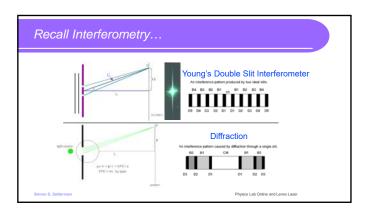
Liquid-Core/ Cladding (L2) Waveguide

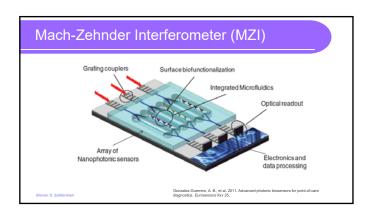


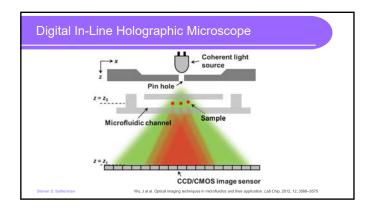
Three parallel waveguides formed with liquid core and cladding in laminar flow systems. The direction of the light propagation can be altered by different flow rates of the adjacent fluids.

Gai, Hongwei, Yongjun Li, and Edward S. Yeung. 2011. Op Technologies and Applications 304, 171-201. Ligler FS (2009) Anal Chem 81:519–526









Summary

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- On-Chip Methods
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 Optical Waveguides
 Surface Plasmon
 Resonance
 Interferometry
 Holography

 Appendix
- Appendix
- Tables of optical detection methods

Table of Chemiluminescence Detection

| Optical Detection | Sensor Technology | Analyte | Assay Type | Time of Analysis | Resolution | Point of Care* | Ref. |
|-------------------|-----------------------------|---------------------------------|--------------------------|---------------------|---------------------|-------------------|------|
| Chemiluminescence | Microplate reader | Hepatitis B antigen | Capillary immunoassay | 25 min | 0.3 ng/mL | 4 | [52] |
| Chemiluminescence | PMT | Carcinoembryonic autigen | Sandwich immunoassay | 85 | 20 pg/ml. | + | [53] |
| Chemiluminescence | CCD camera | Staphylococcal enterotoxin B | Sandwich immunoassay | >60 min | 0.1 ng/mL | + | [54] |
| Chemiluminescence | Inorganic photodiodes | Anti-HRP antibody | HRP-luminol reactions | >60 min | $0.2~\mathrm{amol}$ | ++ | [55] |
| Chemiliminescence | Inorganic photoconductor | Streptavidin | HRP-luminol reactions | Real time | 4.76 nM | ++ | [56] |
| Chemiluminescence | Organic photodiodes | Staphylococcal enterotoxin B | Sandwich immmosssay | 60-70 s | 0.5 ng/mL | +++ | [57] |

Pires, NM, et al. Recent Developments in Optical Detection Technologies in Lab-on-a-Chip Dev Biosensing Applications. Sensors 2014, 14, 15458-15479

| • | 52. Xiang, A.; Wei, F.; Lei, X.; Liu, Y.; Liu, Y.; Guo, Y. A simple and rapid capillary chemilluminescence immunoassay 1 quantitatively detecting human serum HBsAg. Eur. J. Clin. Microbiol. Infect. Dis. 2013, 32, 1557–1564. | lo |
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| | quantitatively detecting numeri serum ribsag. Eur. 3. Olin. Microbiol. Milect. Dis. 2019, 32, 1337-1304. | |

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Table of Fluorescence Detection

| Optical Detection | Sensor Technology | Analyte | Assay Type | Time of Analysis | Resolution | Point of Care* | Ref. |
|-------------------|--------------------------|--------------------------------|----------------------------|---------------------|--------------------------|-------------------|------|
| Fluorescence | CMOS image sensor | Giardia Lamblia cysts | Microscopy | -1 s | Focal plane of 0.8 µm | * | [41] |
| Fluorescence | CCD camera | Bacterial DNA | PCR | Real time | -50 CFU/mL ³ | + | [49] |
| Fluorescence | Inorganic photodiodes | 17-β estradiol | Competitive aptamer assay | ~10 min | 0.6 ng/mL | ++ | [50] |
| Fluorescence | Organic photodiodes | Alkylphenol polyethoxylates | Competitive immunoassay | -5 min | 2-4 ppb | ++ | [51] |

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Pires, NM, et al. Recent Developments in Optical Detection Technologies in Lab-on-a-Chip Devices & Biosensing Applications. Sensors 2014, 14, 15458-15479

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 microchannel. *Talanta* 2013, 117, 139–145.

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