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Phase-sensitive Photoelectron Metrology

Paul Hockett

femtolab.ca National Research Council of Canada, Ottawa **Available via Figshare, DOI: [10.6084/m9.figshare.5049142](http://dx.doi.org/10.6084/m9.figshare.4906889)**

Interferometers...

matter wave interferometry

Young's double slit with electrons... two-path quantum interferometry.

Figure credit: Simon Connell. http://psi.phys.wits.ac.za/teaching/connell/phys284/2005/lecture-02

general quantum interferometry $TATATATOMATIV$ in lonization paths of \sim \sim there by producing a phase shift in the observed signal. An analytical theory was used to determine the observed to determine the observed signal. An analytical theory was used to determine the observed to determine the s

... pretty much any quantum mechanical process where *multiple paths* play a role. <u>The chailed process where *indiuple paths*</u> gredient in all quantum mechanical interference phenom-Fig. 1. As in any interferometric measurement, two phase-

and/or short-range resonance effects. Long-range forces

it is the phase that care the interest For example, the phase of a bound-state wavefunction:

VOLUME 85, NUMBER 10 PHYSICAL REVIEW LETTERS 4 SEPTEMBER 2000 $\frac{4 \text{ J}}{\text{L}}$ phases $\frac{4 \text{ J}}{\text{L}}$ and $\frac{2000}{\text{L}}$

Direct Observation of a Breit-Wigner Phase of a Wave Function ase of a wave function

Jeanette A. Fiss, Ani Khachatrian, Kaspars Truhins, Langchi Zhu, and Robert J. Gordon Department of Chemistry (m/c 111), University of Illinois at Chicago, 845 West Taylor Street, Chicago, Illinois 60607-7061

Tamar Seideman

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Steacie Institute for Molecular Sciences, National Research Council of Canada, Ottawa, Canada K1A-0R6 (Received 16 February 2000) cil of Canada, Ottawa, Canada KIA-0R6
00) T

The Breit-Wigner phase of a wave function was obtained by measuring the interference between two independent ionization paths of a molecule. The state of interest was present in only one of the paths, thereby producing a phase shift in the observed signal. An analytical theory was used to determine the phase of the wave function from the observable.

FIG. 1. Illustration of a molecular interferometer. Panel (a) shows that two competing quantum mechanical paths connecting the same initial and final states produce a sinusoidal variation of the product signal that depends on the relative phase of the two paths. In panel (b) an additional phase source is introduced at an intermediate (two-photon) level of the three-photon path. This source could be, for example, a predissociating resonance. The effect of this source is to produce a phase shift of δ_{13} in the signal.

general quantum interferometry

... pretty much any quantum mechanical process where *multiple paths* play a role.

For example, the phase of the free electron wavefunction(s) via multiple ionization paths:

VOLUME 69. NUMBER 16

PHYSICAL REVIEW LETTERS

19 ОСТОВЕВ 1992

Asymmetric Photoelectron Angular Distributions from Interfering Photoionization Processes

Yi-Yian Yin, Ce Chen, and D. S. Elliott School of Electrical Engineering, Purdue University, West Lafayette, Indiana 47907-1285

A. V. Smith

Sandia National Laboratories, Albuquerque, New Mexico 87185 (Received 14 May 1992)

We have measured asymmetric photoelectron angular distributions for atomic rubidium. Ionization is induced by a one-photon interaction with 280 nm light and by a two-photon interaction with 560 nm light. Interference between the even- and odd-parity free-electron wave functions allows us to control the direction of maximum electron flux by varying the relative phase of the two laser fields.

PACS numbers: 32.80.Fb, 32.80.Rm

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Note - control via E-field phases

FIG. 3. Experimental data. The total electron count as a function of pressure of N_2 gas in the phase delay cell for the four detectors positioned at (a) 0° , (b) 45° , (c) 90° , and (d) 180°. The solid line is the result of a least-squares fit of a sinusoidally varying curve to the data.

general quantum interferometry energy functions from spectroscopic data [8]. The phase of a scattering wave function is conveniently separated into a dynamical and a geometric part by partial $\overline{}$ gredient in all quantum mechanical interference phenomena. These include \overline{C} arol quontu Igrafiuudilu coherent pathways connect the initial and final points.

... pretty much any quantum mechanical process where *multiple paths* play a role. the molecular orientation and on the scattering molecular orientation and on the scattering orientation and or Dynamical phases arise from long-range scattering effects adva loie. scattering wave function as compared with those of a free $[6,7]$. In particular, the phases of continuum wave functions of continuum wave functions of continuum wave functions \mathcal{L} \blacksquare tering dynamics is explicit in the form of the form of the form of the *pic* function and in inversion schemes used to extract potential molecular interferometer, the two pathways consist of relly much any quant ionization continuum (Fig. 1a). The relative phase of the

femtolab.ca (a) from Fiss et. al., PRL 85 2096 2000 *femtolab.ca*

photoionization interferometry

Photoionization is an interferometric process, in which multiple paths can contribute to the final continuum photoelectron state.

(1) Intrinsic: interferences between final (continuum) states.

(2) Extrinsic/dynamic: additional pathways due to, e.g., prepared wavepacket, control fields, etc. etc.

phase-sensitive photoelectron metrology

Need a phase-sensitive observable... photoelectron angular distributions (PADs) are angle-resolved photoelectron interferograms.

Although there are other factors, this illustrates why the PAD is so sensitive to the phase shifts - it is the interference due to these phase shifts which primarily determines the shape of the PAD.

phase-sensitive photoelectron metrology

Need a phase-sensitive observable... photoelectron angular distributions (PADs) are angle-resolved photoelectron interferograms.

s+p(m=0)+d(m=1) waves, as a function of relative phase.

background - dipole matrix elements

Any measurement involving ionization projects the initial state wavefunction onto the ionization continuum - free electron + ion.

In the weak-field & dipole regime, this is described by the dipole matrix elements:

Observable - angle-resolved photoelectron flux: $U($

$$
I(\theta, \phi; E, t) = \langle \Psi_+; \psi_e | \hat{\mu} . \mathbf{E} | \Psi_i \rangle \langle \Psi_i | \hat{\mu} . \mathbf{E} | \Psi_+; \psi_e \rangle
$$

$$
=\sum_{L}\sum_{M}\beta_{LM}(E,t)Y_{LM}(\theta,\phi)
$$

By writing all the wavefunctions as products of radial & angular (geometric) parts, the cross-section can be written as:

$$
I(\theta, \phi; E, t) = \sum_{l\lambda} \sum_{l'\lambda'} \gamma r_{l\lambda} r_{l'\lambda'} \cos(\eta_{l\lambda} - \eta_{l'\lambda'}) Y_{l\lambda}(\theta, \phi) Y_{l'\lambda'}^*(\theta, \phi)
$$

The summation shown here is over all angular momentum states of the free electron, hence the PAD is an *(angular) interference pattern.*

But... there may be many channels involved!

Isn't this just photoelectron spectroscopy?

Isn't this just photoelectron spectroscopy?

Absolutely... but with a focus on (quantitative) phase-sensitive metrology.

Measurement sophistication (information content)

background - radial integrals

The geometric terms are analytic, so can be calculated directly, leaving only the radial integrals as unknowns. How can we treat these?

$$
I(\theta, \phi; E, t) = \sum_{l\lambda} \sum_{l'\lambda'} \gamma r_{l\lambda} r_{l'\lambda'} \cos(\eta_{l\lambda} - \eta_{l'\lambda'}) Y_{l\lambda}(\theta, \phi) Y_{l\lambda}^*(\theta, \phi)
$$

Radial integrals \equiv scattering amplitudes & phases
Ab initio (numerical)
scattering calculation
Quantitative methods
Symmetry based modelling
Qualitye methods

recent examples

Complete Photoionization Experiments via Ultrafast Coherent Control with Polarization Multiplexing

Hockett, P., Wollenhaupt, M., Lux, C., & Baumert, T. Physical Review Letters, 112(22), 223001 (2014). http://doi.org/10.1103/PhysRevLett.112.223001 arXiv 1403.3315 (https://arxiv.org/abs/1403.3315)

Angle-resolved RABBIT: theory and numerics

Paul Hockett *J. Phys. B (under review, 2017), arXiv 1703.08586 (https://arxiv.org/abs/1703.08586). Authorea https://dx.doi.org/10.22541/au.149037518.89916908*

Bootstrapping to the Molecular Frame with Time-domain Photoionization Interferometry

Claude Marceau, Varun Makhija, Dominique Platzer, A. Yu. Naumov, P. B. Corkum, Albert Stolow, D. M. Villeneuve and Paul Hockett *Phys. Rev. Lett. (under review, 2017),*

[arXiv 1701.08432 \(https://arxiv.org/abs/1701.08432\)](https://arxiv.org/abs/1701.08432).

Modelling the dynamics & ionization for this 3-photon process.

าล 0.8 *"Control" via interferences:* Population iwayo L vs R pathways to same final states.

femtolab.ca polarization is described by this decomposition, provided $f\Omega$

Experiments (Christian Lux)

Experiments (Christian Lux)

and identical pulse lengths of 80 fs. All pulse durations given here are FWHM values resulting from 30 fs input pulses. As

Fig. 12 *Upper panels*: measured PADs from excitation with polariza-

femtolab.ca are sharply truncated, and therefore, the corresponding tem-

multi-path atomic interferometry \mathbf{b} is the fine structure splitting of the \mathbf{a} state into 4*p*1*/*2- and 4*p*3*/*2-states of about 7 meV is small compared to the last \mathbf{y} at Half Maximum (FWHM) for a 30 fs last public state public state public state public state public state public s

extracting the ionization dynamics

theory & experiment comparison

dynamics & control

Conclusions & future work

- Modelling as 1-photon Rabi-cycling (*intrapulse dynamics*), followed by 2-photon ionization, seems to be valid.
- 2D images provide a route to **extraction of the ionization matrix elements** ("complete photoionization experiment").
- For potassium we determine a dominant *f*-wave, with significant *p*-wave contribution.
- We can now model the **dynamics of polarization control**.
- *• Fuller treatment of radial distributions to model, e.g., chirped laser pulse.*
- *• 3D data (tomographic measurements) should be more sensitive, and allow more* **ust** *fitting.*

for more...

Coherent Control of Photoelectron Wavepacket Angular Interferograms

Hockett, P., Wollenhaupt, M., Lux, C. & Baumert, T. *J. Phys. B, 48, 214004 (2015)* http://doi.org/10.1088/0953-4075/48/21/214004 arXiv 1505.00035

Maximum Information Photoelectron Metrology

Hockett, P., Lux, C., Wollenhaupt, M. & Baumert, T. *Phys. Rev. A, 92, 013412 (2015)* http://doi.org/10.1103/PhysRevA.92.013412 arXiv 1503.08308

Complete Photoionization Experiments via Ultrafast Coherent Control with Polarization Multiplexing II: Numerics & Analysis Methodologies

Hockett, P., Wollenhaupt, M., Lux, C. & Baumert, T. *Phys. Rev. A, 92, 013411 (2015)* http://doi.org/10.1103/PhysRevA.92.013411 arXiv 1503.08247

Complete Photoionization Experiments via Ultrafast Coherent Control with Polarization Multiplexing

P. Hockett, M. Wollenhaupt, C. Lux, T. Baumert *Physical Review Letters 112, 223001 (2014).* http://doi.org/10.1103/PhysRevLett.112.223001 arXiv 1403.3315

recent examples

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Appl. Phys. B 74 [Suppl.], S17–S21 (2002)

DOI: 10.1007/s00340-002-0894-8

Lasers and Optics Applied Physics B

h.g. muller **Reconstruction of attosecond harmonic beating by interference of two-photon transitions**

FOM-Institute for Atomic and Molecular Physics, Kruislaan 407, 1098 SJ Amsterdam, The Netherlands

Received: 19 September 2001/ **Revised version: 7 November 2001 Published online: 5 July 2002 • © Springer-Verlag 2002**

ABSTRACT A method is proposed for detailed determination of the temporal structure of XUV pulses. The method is especially suited for diagnostics on attosecond pulses and pulse trains that originate from temporal beating of various harmonics of an ultrashort laser pulse. A recent experiment already showed the feasibility of this method when applied to long attosecond pulse trains, where it measured the average pulse characteristics. Here we argue that the same method is also suitable for determining differences between the individual attosecond pulses in a short train, or the properties of a single attosecond pulse.

PACS 32.80.Rn; 42.30.Rx

and leave the other pulse unchanged, since the light required for up-conversion by sum-frequency mixing would map in the far or mid infrared. It is much preferable to up-convert both pulses by mixing with an optical photon of slightly different, precisely defined frequency. A convenient way to obtain the narrow-band photons required for the up-conversion process is by selecting different portions of a strongly chirped version of the pulse under study. This pulse is guaranteed to have enough bandwidth to generate a suitable spectral shear Ω.

The major limitation of SPIDER is that to measure pulses that are far from their bandwidth limit, the spectral sampling has to be rather dense (i.e. small Ω). To make a frequency that is sufficiently constant over the (long) duration of the original pulse, one has to chirp out the up-converting pulse so far that it might not have enough intensity left. In cases like this, however, only a small spectral shear is desired, so that is desired, so that is desired, so that is

1 Measuring spectral phases a stronger, narrower-band source, such as the pump laser of http://doi.org/10.1007/s00340-002-0894-8 Muller, H. G. (2002). Reconstruction of attosecond harmonic beating by interference of two-photon transitions. Applied Physics B: Lasers and Optics, 74, 17–21.

Figure 4. Pump–probe schemes. (a) Traditional pump–probe experiment with two pulses separated in time by τ . (b) Simultaneous pump–probe experiment between a SAP and a few-cycle IR field. (c) Simultaneous pump–probe experiment between an APT and a monochromatic IR field. The narrow purple area represents the attosecond XUV pulse envelope and the broader red area represents the one of the probing laser pulse, while the dotted red lines indicate the corresponding *E*-field.

Figure 5. RABITT method. (a) Photoelectron spectrogram over photon energy and delay between the APT and the IR field. The offset in the modulation of the SBs contains information about the attosecond pulses and the ionization process. (b) Schematic energy diagram over the q dantum paths reading to the same mial energy in SD zq . The experimodulation of the SBs contains information about the attosecond pulses and the ionization process. (b) Schematic energy diagram over the
quantum paths leading to the same final energy in SB 2*q*. The experimental data were quantum paths leading to the same final energy in SB 2*q*. The experimental data were gathered from [56].

the attornation devices bahlström. J. M., L'Huillier, A., & Maguet, A. (2012). Introduction information about attosecond photoionization indirectly \overline{a} in photoionization photoelectron photoelectr $\frac{1}{2}$ Physics, 45(18), 183001. http://doi.org/10.1088/0953-4075/45/18 t_{in} lournal of Dhysics $P: \Lambda$ tomic M ologuk α be and the delay can be controlled with the delay can be controlled with α Darnstrom, J. M., Erfunner, A., & Maquet, A. (2012). Introduction to attosecond
delays in photoionization. Journal of Physics B: Atomic, Molecular and Optical ω , ω , ω , to ω is the electron is not complete the electron is not complete The advantages of the periodic time-structure of APT are periodic time-structure o h., & iviaquet, A. (2012). Introduction i potential from the internal from the internal from the internal co-workers found that the internal co-workers found that the internal co-workers found that the internal co-worker found that the internal co-workers found t Physics, 45(18), 183001. http://doi.org/10.1088/0953-4075/45/18/183001 rather practical: first, it is less demanding experimentally to Dahlström, J. M., L'Huillier, A., & Maquet, A. (2012). Introduction to attosecond

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Femtolab.ca $\rho_{\rm cool}$ and $\rho_{\rm CMB}$ and $\rho_{\rm CO}$ for streaking, thereby, causing less side-effects on the system, \mathbf{r} essuur **politicis temmorad** c the signal is read out on zero-background, i.e. in energetic

RABBIT in detail

Each channel contains multiple paths and phases.

Channels: $s \xrightarrow{ xuv } p \xrightarrow{ir } s+d$.

To obtain maximum information, angular resolution is preferable.

m o d e lling AR-RABBIT

Channels: $s \stackrel{xuv}{\rightarrow} p \stackrel{ir}{\rightarrow} s+d$.

To obtain maximum information, angular resolution is preferable.

Realistic numerical model:

- Bound-free (xuv) matrix elements from ePolyScat. [http://www.chem.tamu.edu/rgroup/lucchese/ePolyScat.E3.manual/](http://www.chem.tamu.edu/rgroup/lucchese/ePolyScat.E3.manual/manual.html) [manual.html](http://www.chem.tamu.edu/rgroup/lucchese/ePolyScat.E3.manual/manual.html)
- Continuum-continuum (ir) matrix elements using Coulomb scattering solutions (cf. treatment by Dahlström et. al).

Dahlström, J. M., L'Huillier, A., & Maquet, A. (2012). Introduction to attosecond delays in photoionization. Journal of Physics B: Atomic, Molecular and Optical Physics, 45(18), 183001. http://doi.org/10.1088/0953-4075/45/18/183001

numerical results

AR-RABBIT bands (odd + even harmonics), $I(\theta,\tau)$ **t0.1fs t0.8fs t1.5fs t2.2fs E**

VMI simulation

AR-RABBIT VMI simulation, and comparison with experiments (2 bands).

femtolab.ca Experimental results - Hiromichi Niikura (*Science*, in press, 2017) Model results also include calculated XUV phases - David Villeneuve, SFA calculations

RC-CNRC

AR-RABBIT is a little bit complex, but is an information rich measurement...

It is suitable for both pulse and photoelectron metrology, and control.

Numerical modelling using established photoionization techniques reproduces the expected phenomena, and yields detailed understanding.

The numerical modelling techniques are general (any atom or molecule).*

(Preliminary) comparisons with experimental results (Hiromichi Niikura**) look promising...

* although the method used here is expected to be poor at low (near threshold) photoelectron energies due to the form of the continuum-continuum functions. Strong field effects are neglected.

** *Coherent Imaging of an Attosecond Electron Wave Packet*, D M Villeneuve, P Hockett, M J J Vrakking and H Niikura, Science (in press, 2017)

AR-RABBIT for Wigner delays

For molecules the continuum phase, hence Wigner (photoionization) delay, is a complex function of energy and angle.

AR-RABBIT is one potential method for mapping this phase-dependence.

See:

Time delay in molecular photoionization

P Hockett, E Frumker, D M Villeneuve and P B Corkum, *J. Phys. B: At. Mol. Opt. Phys. 49, 095602, 2016.*

http://dx.doi.org/10.1088/0953-4075/49/9/095602 arXiv 1512.03788

recent examples

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Experimental PADs combined with detailed analysis & theory offer the potential for a move beyond current phenomenological time-resolved imaging techniques by utilizing the photoelectron interferometer.

Experimental PADs combined with detailed analysis & theory offer the potential for a move beyond current phenomenological time-resolved imaging techniques by utilizing the photoelectron interferometer.

The examples discussed so far show some of this potential...

To proceed, we can consider "maximum information" experimental measurements, which allow for determination of the partial waves as a function of time.

One example is the use of impulsive alignment techniques (rotational wavepackets).

Align-probe angleresolved measurements from N_2 (hv=23.3eV).

rotational wavepacket interferometry

Fitting such data as a function of alignment can provide the ionization matrix elements and phases.

Essentially, the rotational wavepacket acts as a geometric contribution to the interferometer.

molecular frame reconstruction

For N_2 , this has been demonstrated for matrix element retrieval for three different final ion states, and verified via molecular frame reconstruction & comparison with theory (ePolyScat).

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[arXiv 1701.08432 \(https://arxiv.org/abs/1701.08432\).](https://arxiv.org/abs/1701.08432)

quantitative molecular dynamics

Most generally, we can look at the full $\beta_{LM}(E,t)$ spectra...

Full excited state molecular dynamics & observable c alculations for CS₂.

quantitative molecular dynamics

Most generally, we can look at the full $\beta_{LM}(E,t)$ spectra...

... and the underlying partial wave amplitudes and phases.

Phase

Full excited state molecular dynamics & observable c alculations for CS₂.

quantitative molecular dynamics

Most generally, we can look at the full $\beta_{LM}(E,t)$ spectra...

... and the underlying partial wave amplitudes and phases.

 100

-50

 θ

 10

 -10

Phase/red.

íο

100

100

200

200

300

300

400

t/fs

400

t/fs

500

500

600

600

700

700

800

Other examples of "maximum information" measurements include tomographic imaging, multi-path ionization schemes and complex light-matter interactions.

Quantum dynamical imaging

The tools are now in place, we just need to use them!

Quantum Dynamical Imaging via Time-resolved Photoelectron Interferometry: Beyond a Phenomenological Imaging of Molecular Dynamics P. Hockett (research proposal, 2013) Available on Figshare, https://dx.doi.org/10.6084/m9.figshare.3580734

a c k n o w l e d g e m e n t s

NRC

Claude Marceau Varun Makhija Ruaridh Forbes Rune Lausten Albert Stolow Eugene Frumke David Villeneuve Paul Corkum Michael Schuurman

& everyone else at NRC!

Caltech Kwangshi Wang Vince McKoy

Texas A&M Robert R. Lucchese (ePolyScat)

Kassel University, Germany

Christian Lux Matthias Wollenhaupt Thomas Baumert

University of Nottingham, UK Katharine Reid

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Web: femtolab.ca

We're always interested in new collaborations and new directions...

If you have an idea, or work that could benefit from our expertise and facilities...

...*please get in touch!*

Slides available via Figshare, DOI: 10.6084/m9.figshare.5049142

Web: www.femtolab.ca

Coming soon: **Quantum Metrology with Photoelectrons [working title]** *New book for the IOP Concise series, due 2018*

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