How to collect best MX data

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CCP4 Workshop, DLS, 2020-11-30

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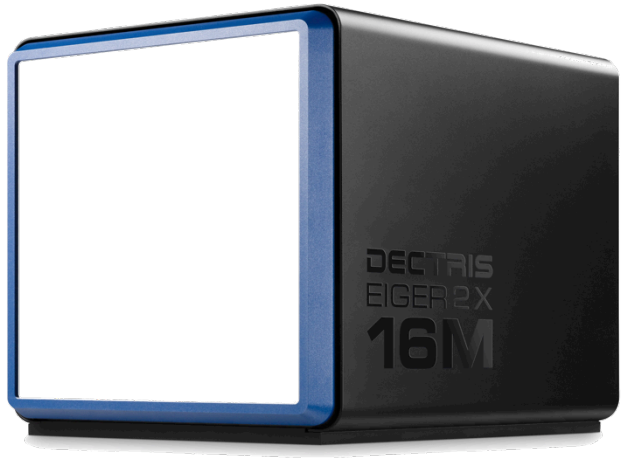
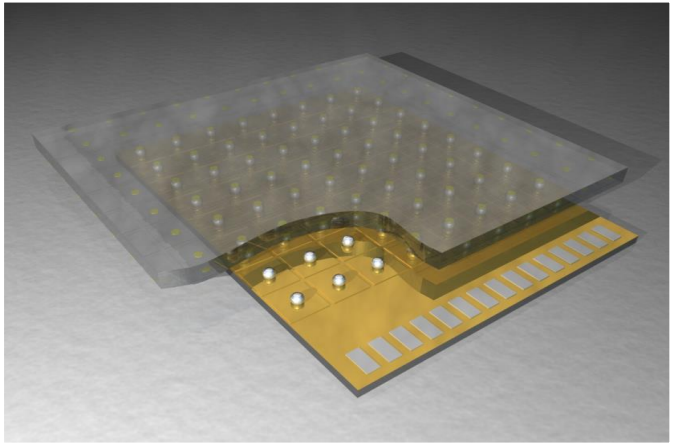
Outline

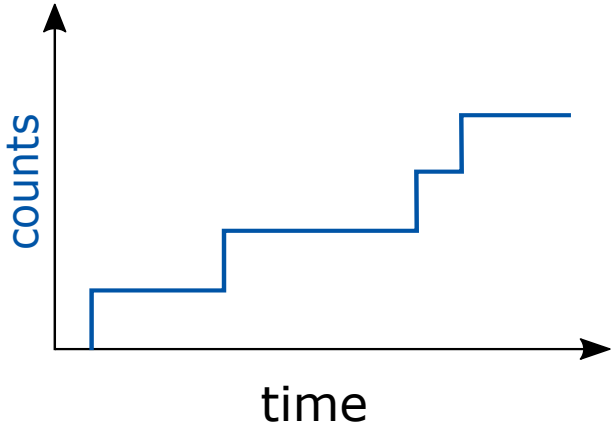
- *Characteristics of X-ray detectors*
- *Accurate data – best practice*
  - *Fine φ-slicing*
  - *Low intensity*
- *Think about your experiment*



The meaning of HPC

Hybrid Photon Counting = Hybrid pixels + photon counting





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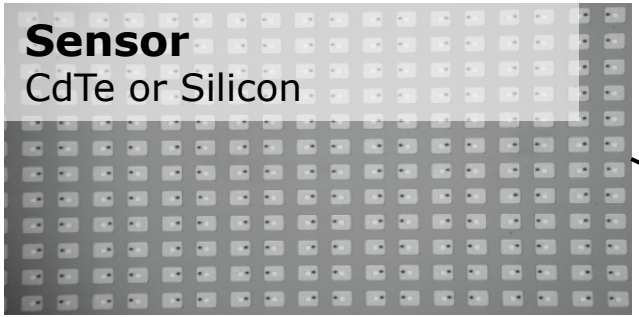
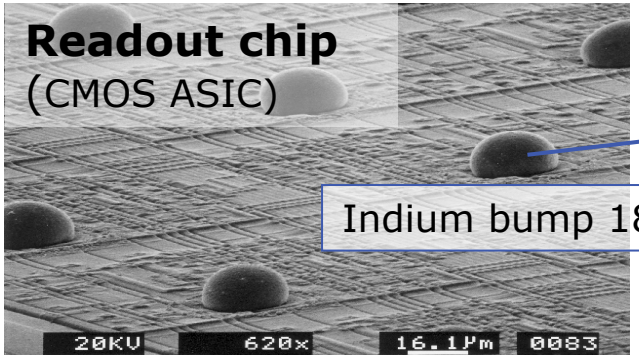
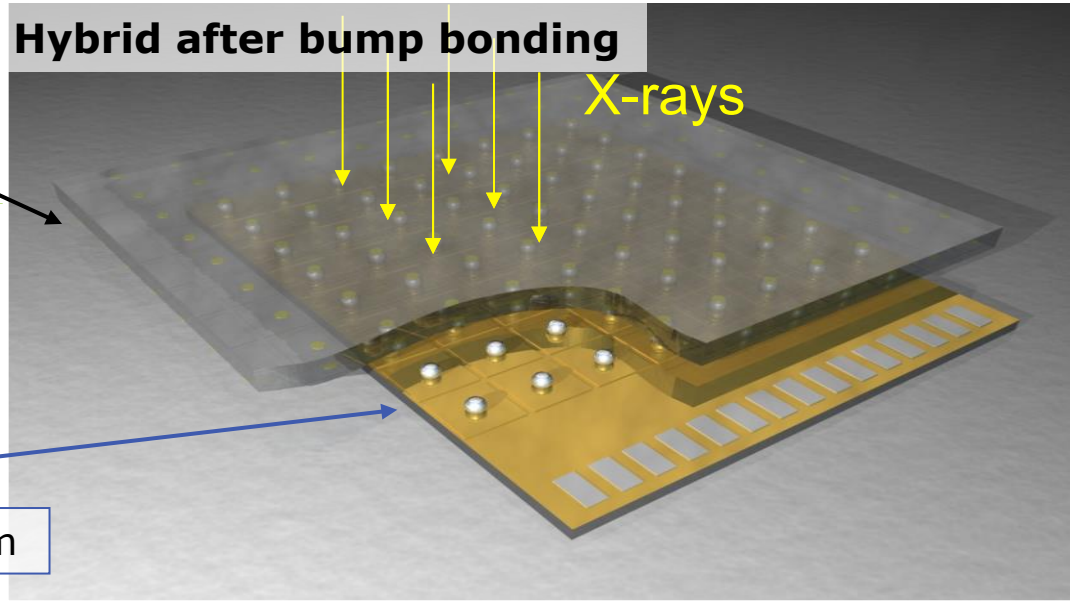
Hybrid pixel = sensor + readout







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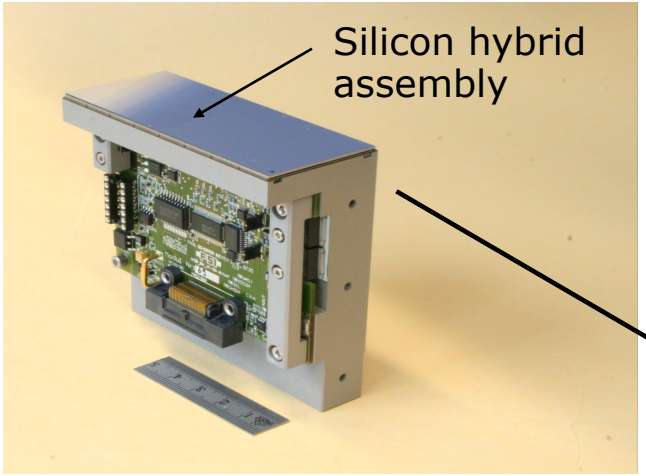
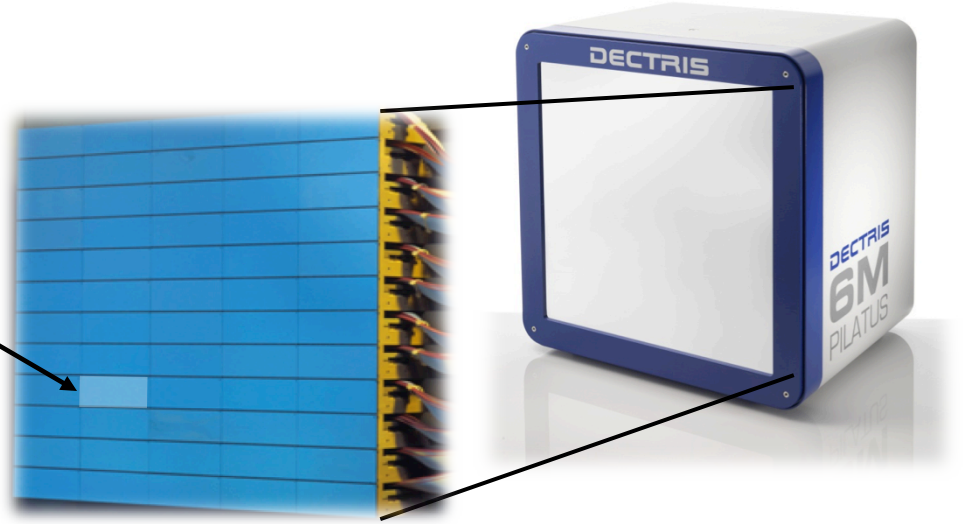
Modular HPC detectors





Module active area: 8 × 4 cm²

100k pixels on PILATUS (172 µm)

500k pixels on EIGER (75 µm)

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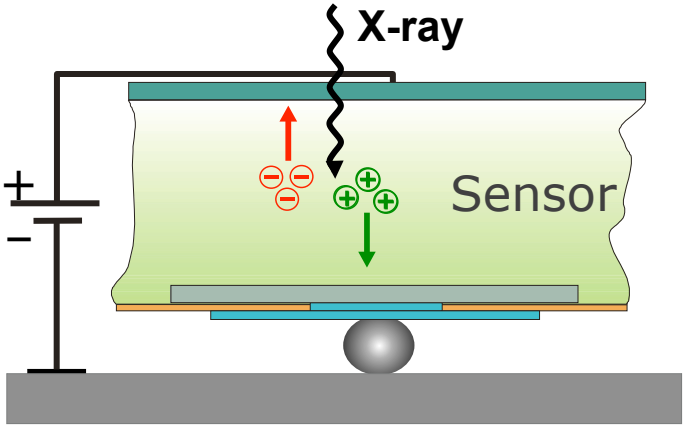
Photon detection in hybrid pixels



Sensor pixel

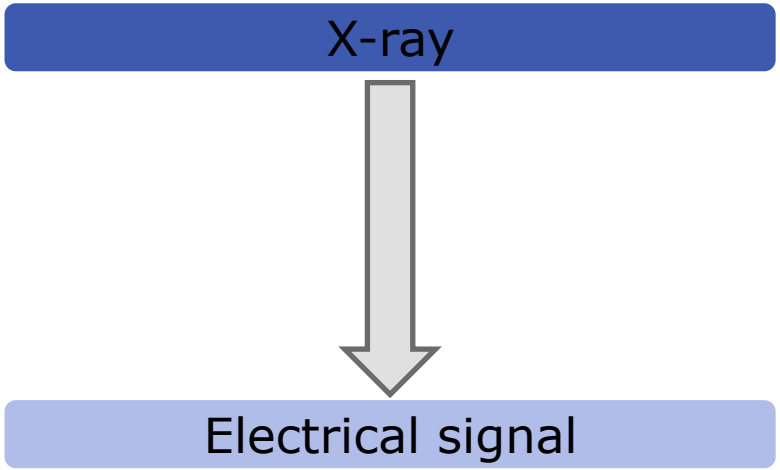
- *Direct detection of X-ray photons -> one e^-/hole pair per 3.6 eV*
- *Charge is captured by electric field*

Readout electronics

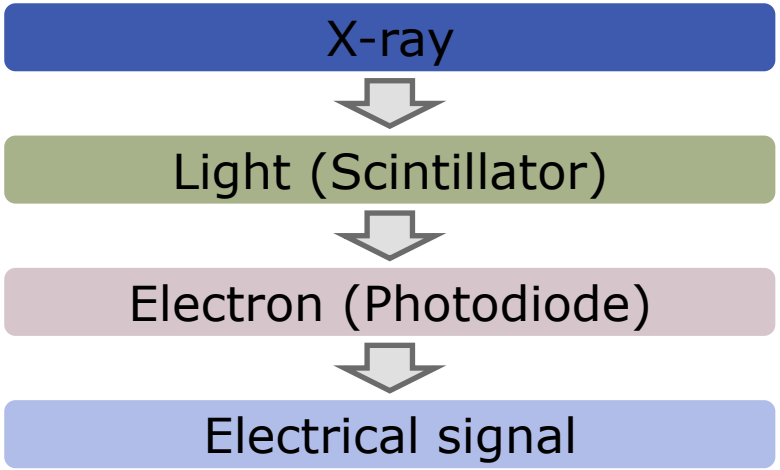
- *Counting of charge pulses*

Superiority of direct detection

Direct detection



Indirect detection



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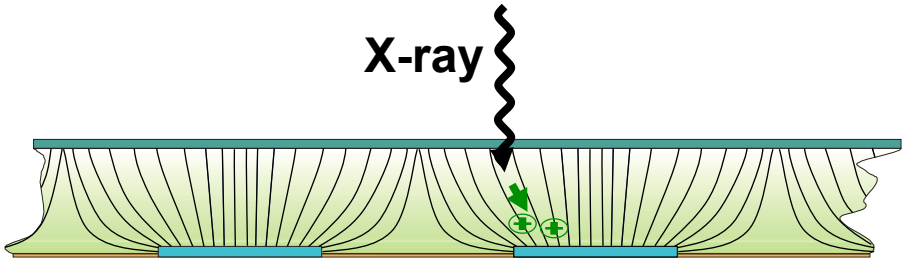
Superiority of direct detection

Direct detection

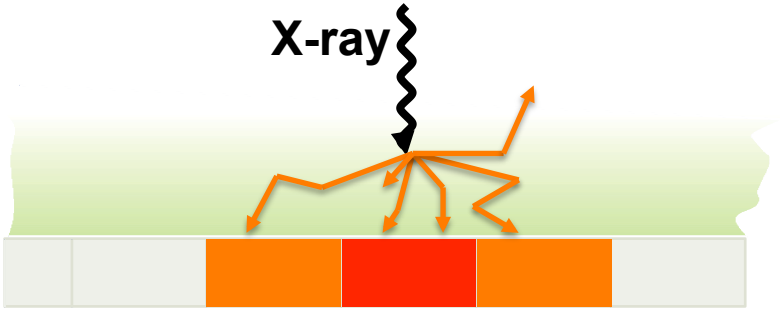


– *Charge captured in electric field*

=> ***All photons captured***

=> ***Signal remains in pixel***

Indirect detection



– *Radiation scattered in scintillator*

=> ***Signal spread across pixels***

=> ***Light partially lost***

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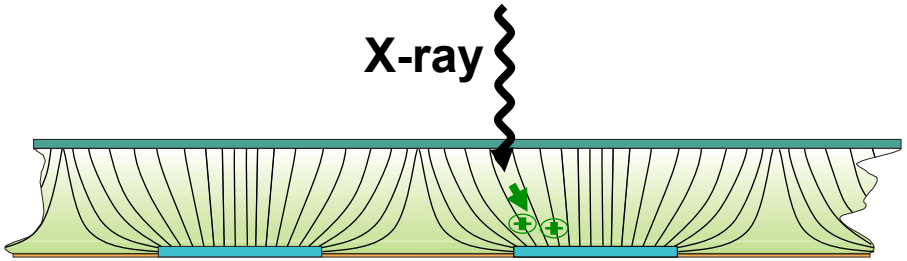
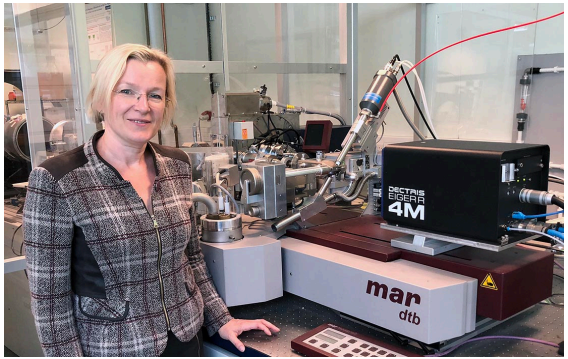
Superiority of direct detection

Charge captured in electric field





- *No photon loss*
- *Sharpest reflections*

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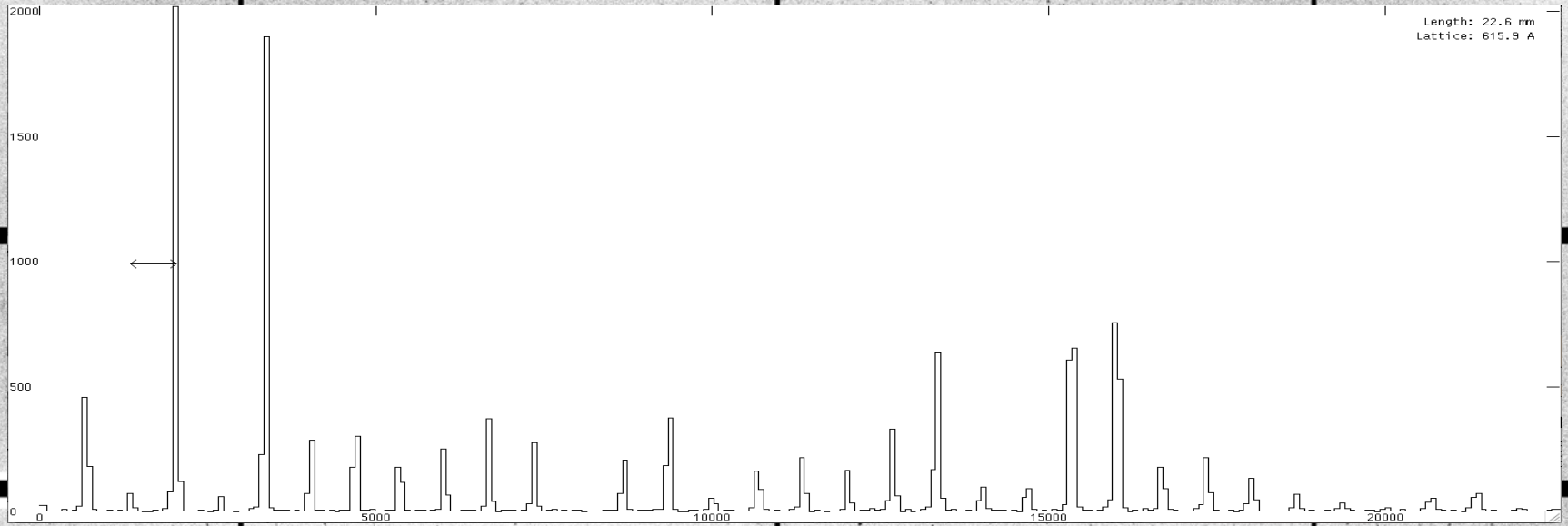
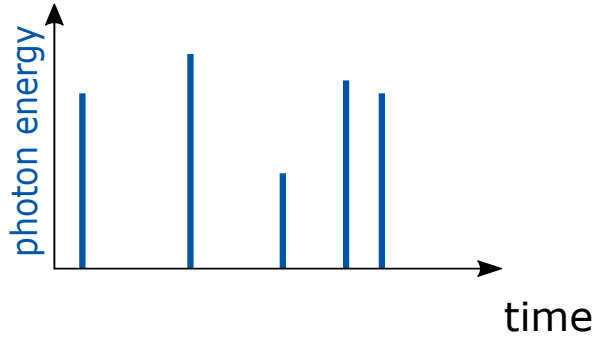
70S ribosome on EIGER X 16M



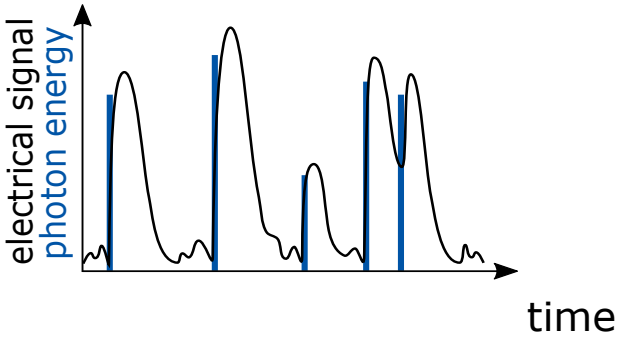
a = 210 Å, b = 450 Å, c = 620 Å
Diffraction to 2.3 Å, Y. Polikanov, UIC

Photon counting – one by one

– *Photons absorbed in sensor pixel*

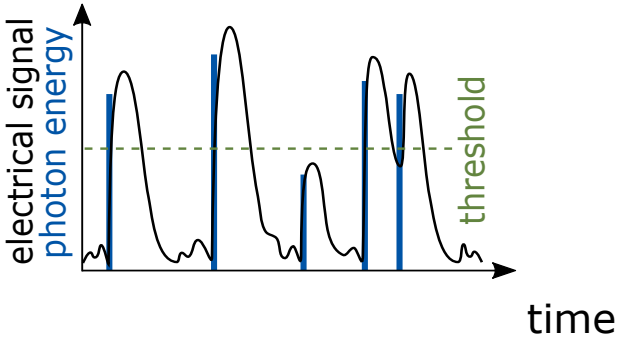


Photon counting – one by one



- *Photons absorbed in sensor pixel*
- *Charge pulse proportional to energy*

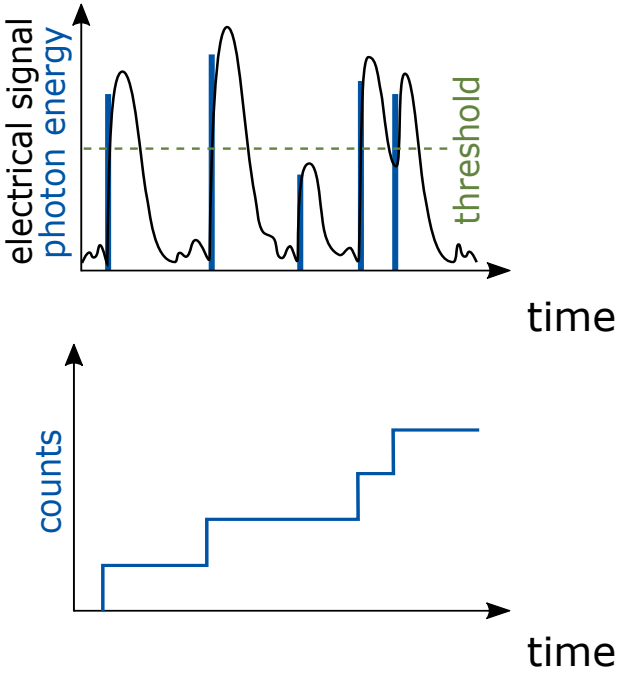
Photon counting – one by one



- *Photons absorbed in sensor pixel*
- *Charge pulse proportional to energy*
- *Threshold to discard noise*

- *Signals above threshold are counted*
  - ***Suppression of dark signal***
  - *Suppression of electronic noise*

Photon counting – one by one



- *Photons absorbed in sensor pixel*
- *Charge pulse proportional to energy*
- *Threshold to discard noise*

- *Signals above threshold are counted*
  - ***Suppression of dark signal***
  - *Suppression of electronic noise*

- *On-the-fly digitization in digital counter*
  - ***No readout noise***
  - *Fast readout*
  - *High dynamic range*

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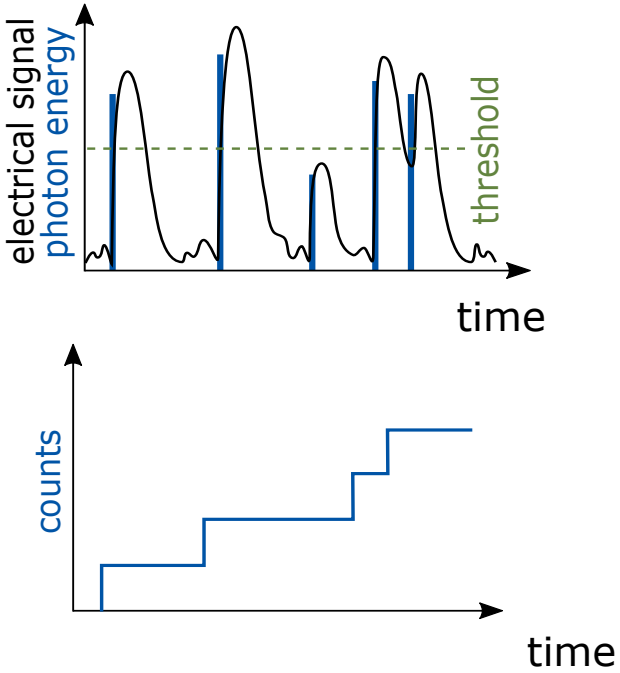
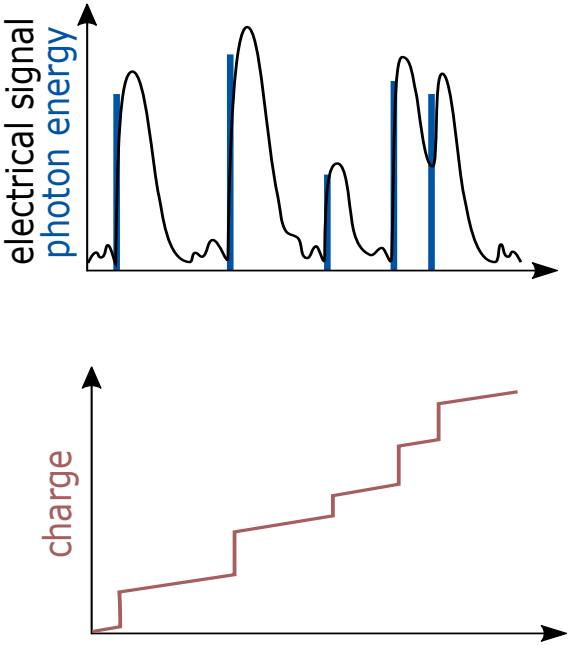
Photon counting vs. integration

Photon counting



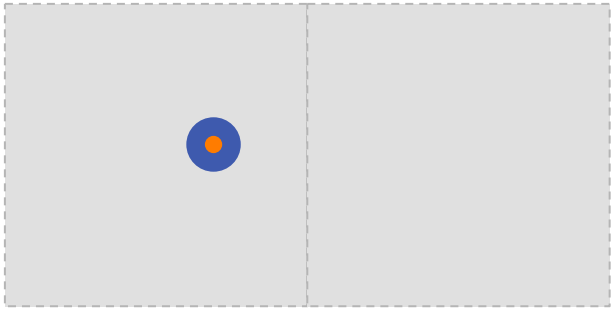
Charge integration



Counting with 50% threshold

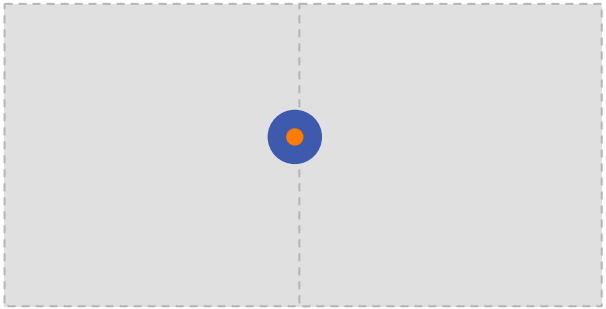
All signal counted within one pixel

1 photon



100% charge
1 count

1 photon



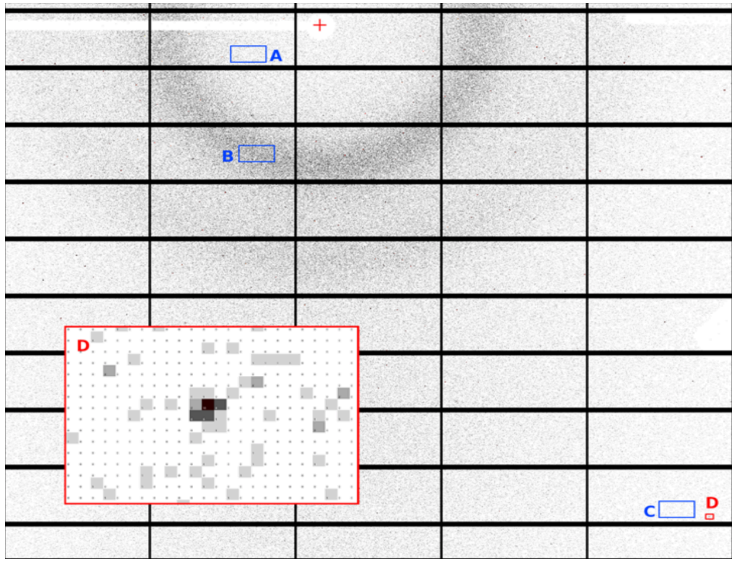
60% charge
1 count

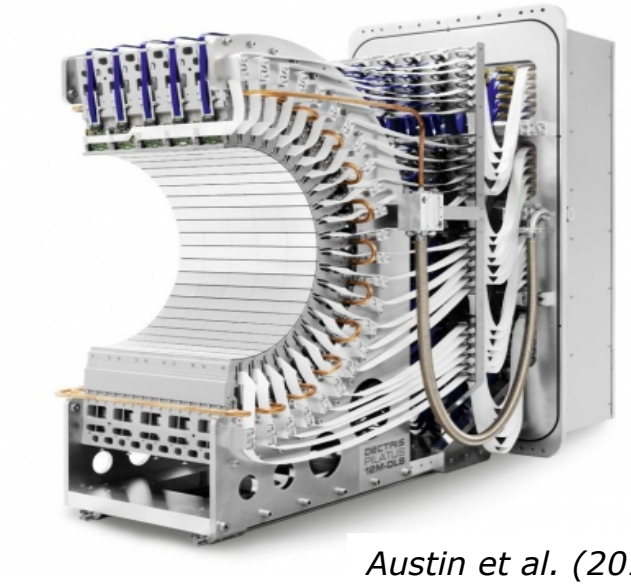
40% charge
-

Negligible background

Protein crystallography in vacuum

PETase @ I23.





Austin et al. (2018)

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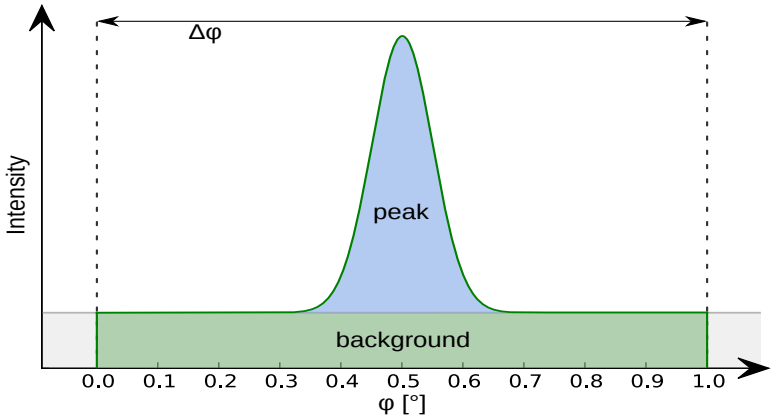
Accuracy of intensity estimates



Q : Number of photons

$var(Q) = Q$

$\sigma(Q) = [var(Q)]^{1/2} = Q^{1/2}$

$I = \Sigma(Peak_i - Bkg_i)$

$var(I) = \Sigma[var(Peak_i) + var(Bkg_i)]$

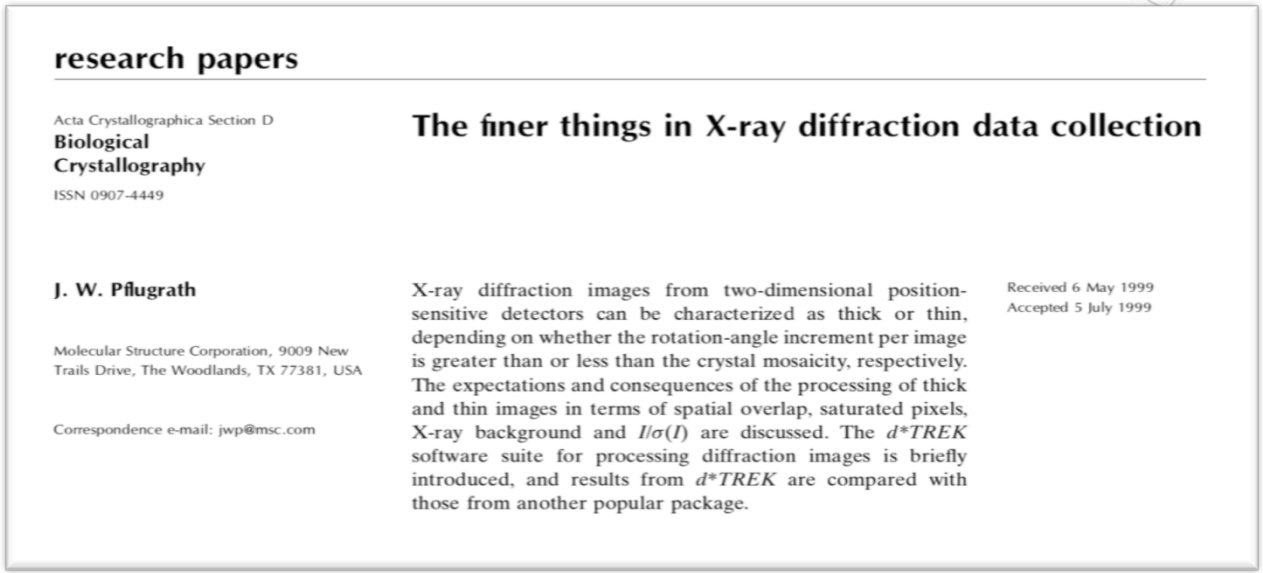
$R_{err} = \sigma(I)/I$

Less background

=> Better data

Relative error [%]	Background [photons]		
Integrated intensity [photons]	0	500	1000
100	10.0	33.2	45.8
500	4.5	7.7	10.0
1000	3.2	4.5	5.5
10000	1.0	1.0	1.1

Pros and cons of fine φ-slicing



research papers

Acta Crystallographica Section D
Biological Crystallography
ISSN 0907-4449

The finer things in X-ray diffraction data collection

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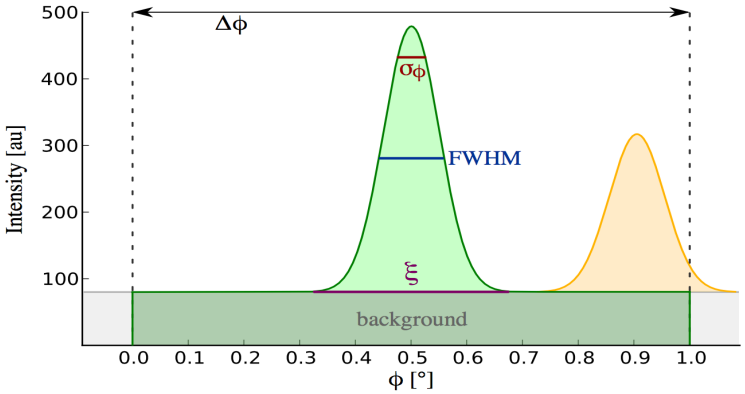
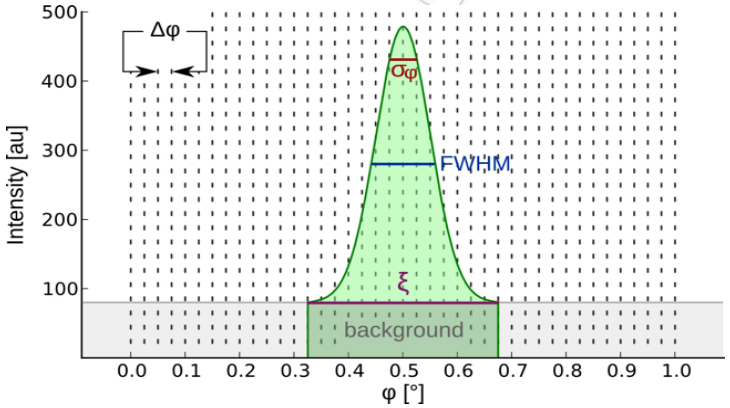
X-ray diffraction images from two-dimensional position-sensitive detectors can be characterized as thick or thin, depending on whether the rotation-angle increment per image is greater than or less than the crystal mosaicity, respectively. The expectations and consequences of the processing of thick and thin images in terms of spatial overlap, saturated pixels, X-ray background and $I/\sigma(I)$ are discussed. The *d*TREK* software suite for processing diffraction images is briefly introduced, and results from *d*TREK* are compared with those from another popular package.

Received 6 May 1999
Accepted 5 July 1999

doi.org/10.1107/S090744499900935X

Fine φ-slicing minimizes background





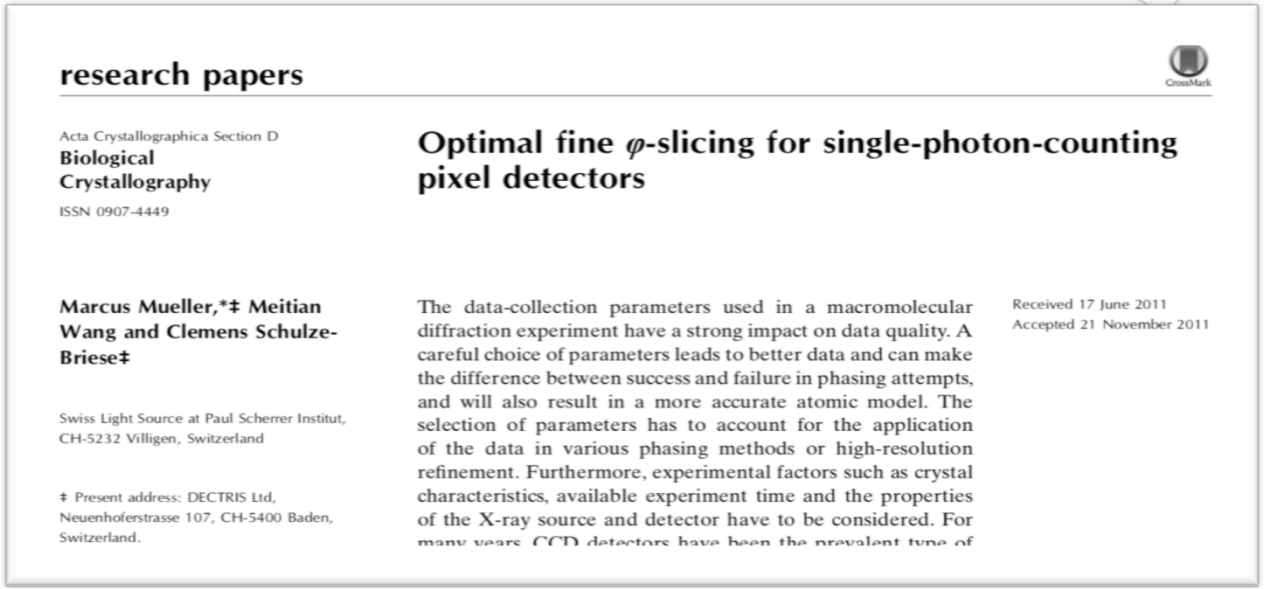
Wide φ-slicing

- Δφ > ξ
- Lots of background
- Few images

Fine φ-slicing

- Δφ ≪ ξ
- Minimal background
- Many images

Advantages of fine φ-slicing



research papers

Acta Crystallographica Section D
Biological Crystallography
ISSN 0907-4449

Optimal fine φ-slicing for single-photon-counting pixel detectors

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The data-collection parameters used in a macromolecular diffraction experiment have a strong impact on data quality. A careful choice of parameters leads to better data and can make the difference between success and failure in phasing attempts, and will also result in a more accurate atomic model. The selection of parameters has to account for the application of the data in various phasing methods or high-resolution refinement. Furthermore, experimental factors such as crystal characteristics, available experiment time and the properties of the X-ray source and detector have to be considered. For many years, CCD detectors have been the prevalent type of

Received 17 June 2011
Accepted 21 November 2011

doi.org/10.1107/S0907444911049833

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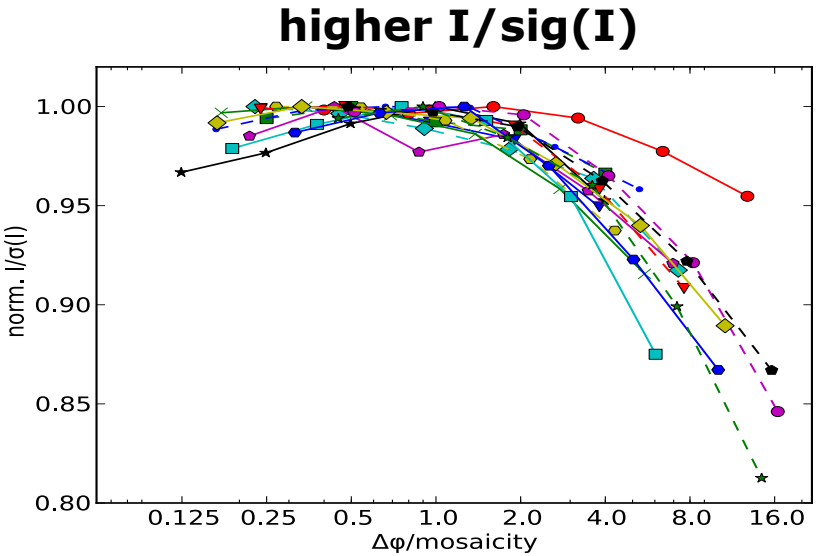
Fine φ-slicing improves data quality

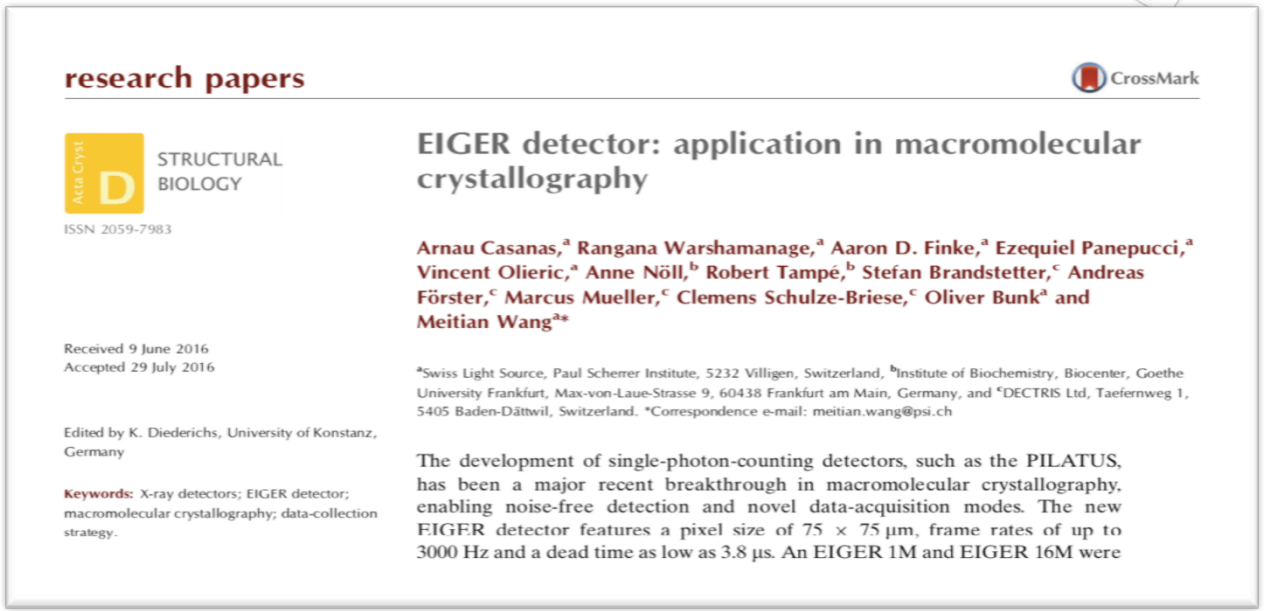
Δφ < mosaicity improves:

- *overall statistics*
- *anomalous signal*
- *highest-shell statistics*
- *number of overlaps*

on PILATUS.



Fine φ-slicing with EIGER



research papers

CrossMark

STRUCTURAL BIOLOGY

ISSN 2059-7983

EIGER detector: application in macromolecular crystallography

Arnau Casanas,[a] Rangana Warshamanage,[a] Aaron D. Finke,[a] Ezequiel Panepucci,[a] Vincent Olieric,[a] Anne Nöll,[b] Robert Tampé,[b] Stefan Brandstetter,[c] Andreas Förster,[c] Marcus Mueller,[c] Clemens Schulze-Briese,[c] Oliver Bunk[a] and Meitian Wang[a]*

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Received 9 June 2016
Accepted 29 July 2016

Edited by K. Diederichs, University of Konstanz, Germany

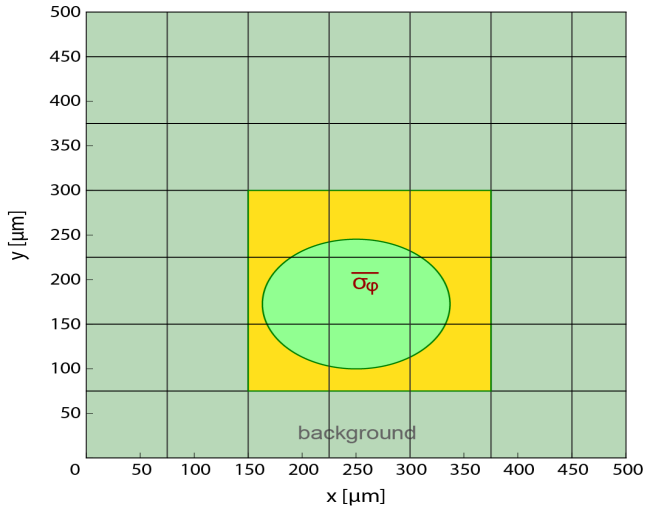
Keywords: X-ray detectors; EIGER detector; macromolecular crystallography; data-collection strategy.

The development of single-photon-counting detectors, such as the PILATUS, has been a major recent breakthrough in macromolecular crystallography, enabling noise-free detection and novel data-acquisition modes. The new EIGER detector features a pixel size of 75 × 75 µm, frame rates of up to 3000 Hz and a dead time as low as 3.8 µs. An EIGER 1M and EIGER 16M were

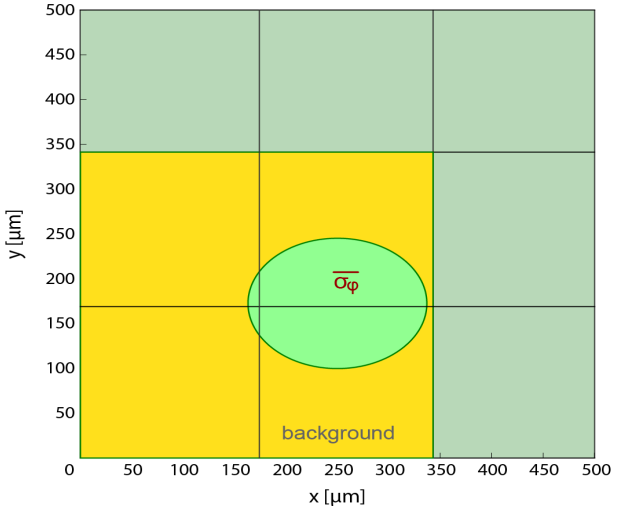
doi.org/10.1107/S2059798316012304

DECTRIS detecting the future

Smaller pixels improve data quality



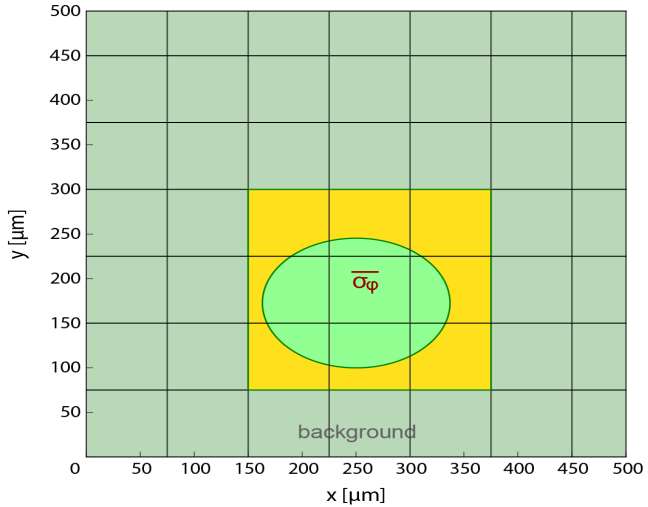
Reflection on EIGER



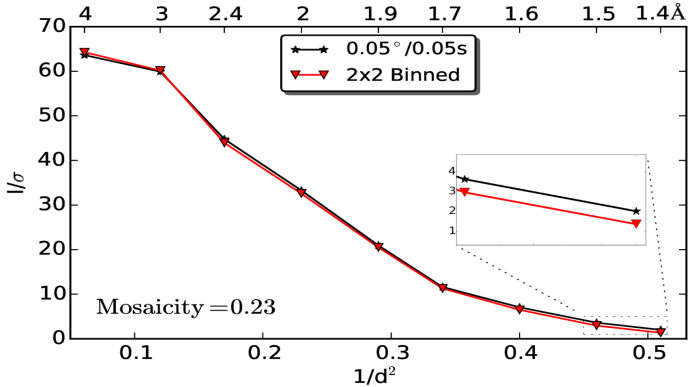
Reflection on PILATUS

DECTRIS detecting the future

Smaller pixels improve data quality



Reflection on EIGER



Better high-resolution data

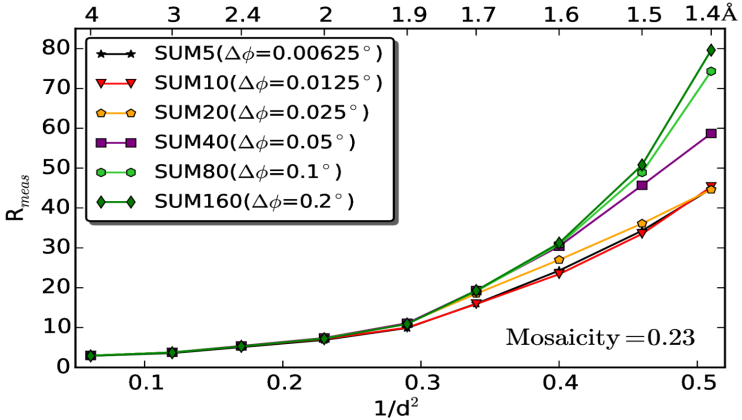
Smaller pixels allow for finer φ-slicing

Δφ ≈ 1/10 mosaicity improves:

- *overall statistics*
- *highest-shell statistics*
- *anomalous signal*
- *number of overlaps*

on EIGER.

lower R_{meas}



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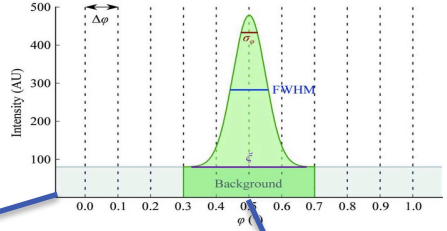
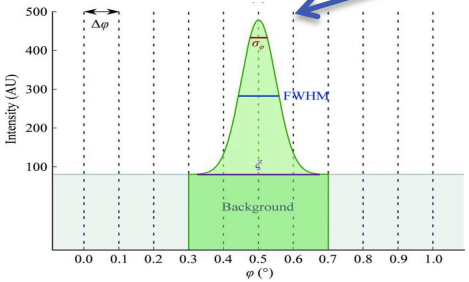
Get best data by

- *Using EIGER and PILATUS detectors*
- *Using fine φ-slicing*
- *Minimizing other sources of error*

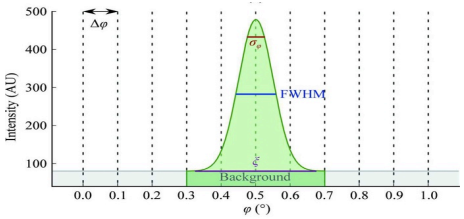
Decrease absolute background



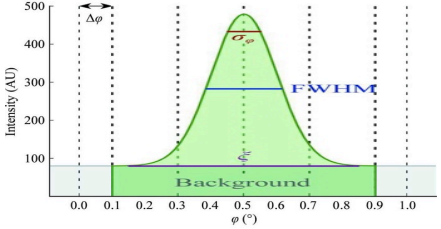
bad!

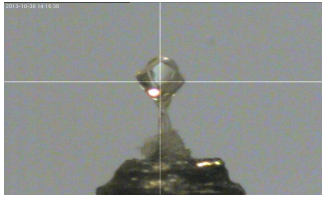


good!



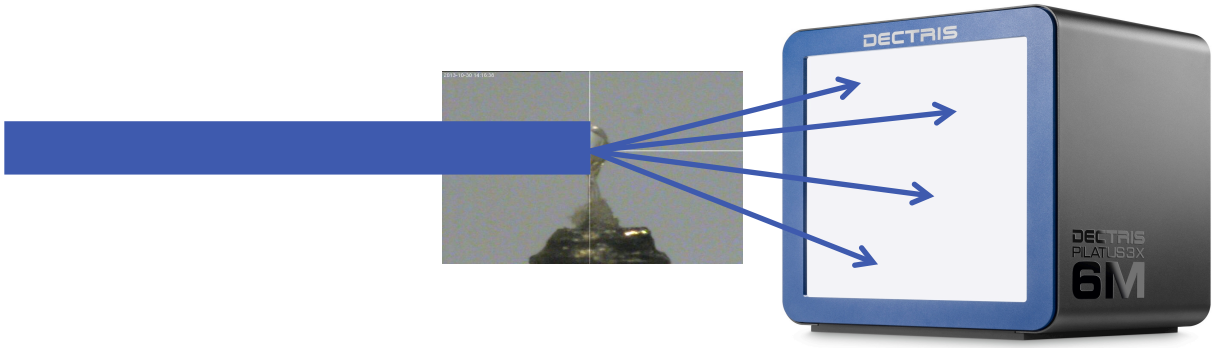
bad!





DECTRIS detecting the future

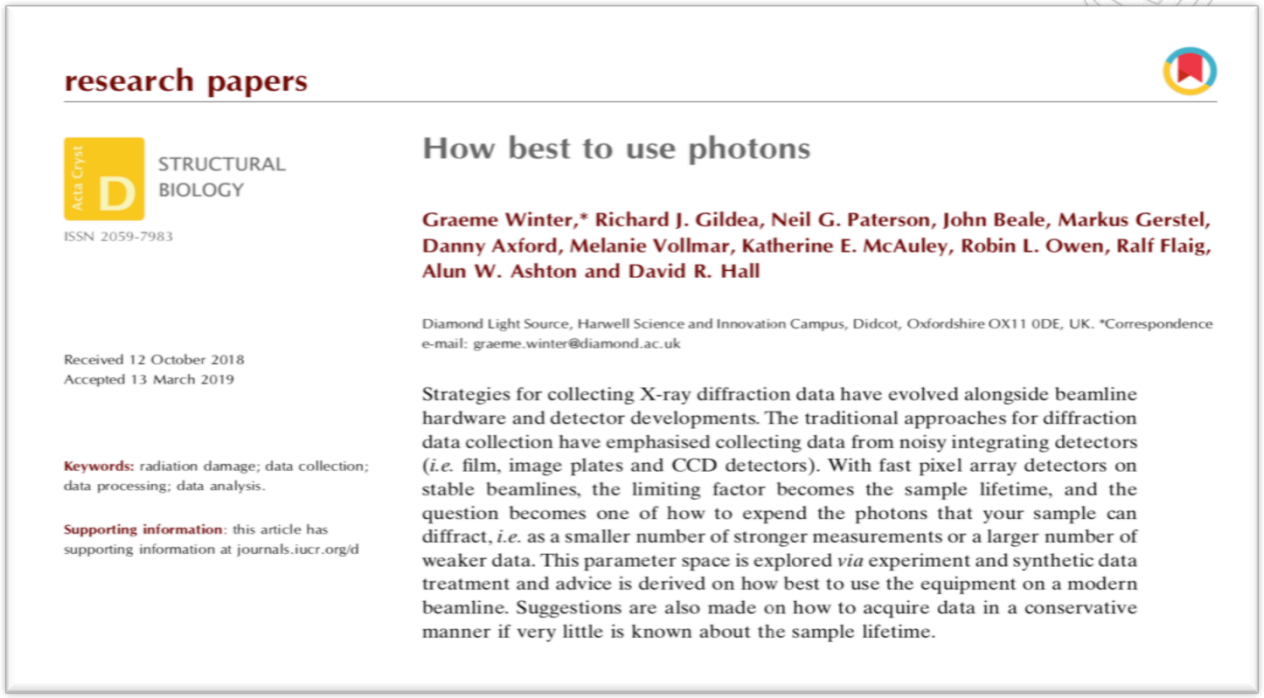
Decrease absolute background



Get best data by

- *Using EIGER and PILATUS detectors*
- *Using fine φ-slicing*
- *Minimizing absolute background*
- *Collecting 360° of data*

Low-intensity data collection



research papers

STRUCTURAL BIOLOGY

ISSN 2059-7983

Received 12 October 2018
Accepted 13 March 2019

Keywords: radiation damage; data collection; data processing; data analysis.

Supporting information: this article has supporting information at journals.iucr.org/d

How best to use photons

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Strategies for collecting X-ray diffraction data have evolved alongside beamline hardware and detector developments. The traditional approaches for diffraction data collection have emphasised collecting data from noisy integrating detectors (*i.e.* film, image plates and CCD detectors). With fast pixel array detectors on stable beamlines, the limiting factor becomes the sample lifetime, and the question becomes one of how to expend the photons that your sample can diffract, *i.e.* as a smaller number of stronger measurements or a larger number of weaker data. This parameter space is explored *via* experiment and synthetic data treatment and advice is derived on how best to use the equipment on a modern beamline. Suggestions are also made on how to acquire data in a conservative manner if very little is known about the sample lifetime.

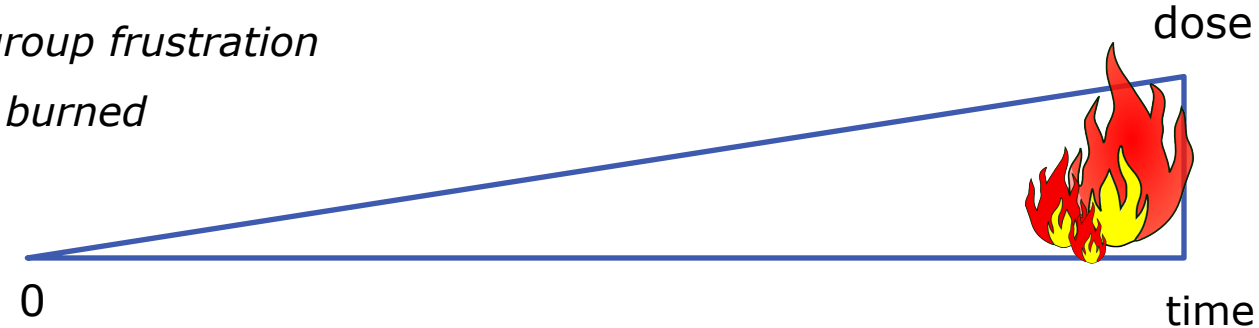
doi.org/10.1107/S2059798319003528

DECTRIS detecting the future

The American method

Collect 360° of data

- *To get more precise intensity estimates*
- *To avoid space group frustration*
- *To avoid getting burned*



	90° @ 10 img/s (0.1°/img)	90 s
	360° @ 40 img/s (0.1°/img)	90 s
Attenuate beam 4-fold +	360° @ 40 img/s (0.1°/img)	90 s

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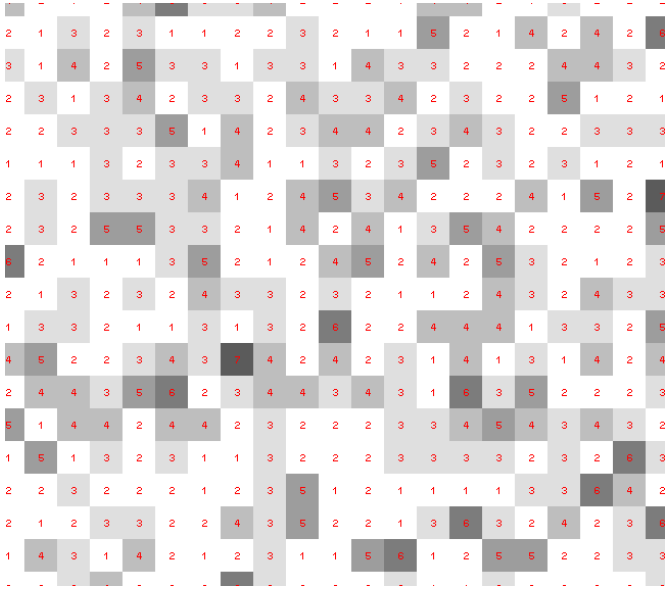
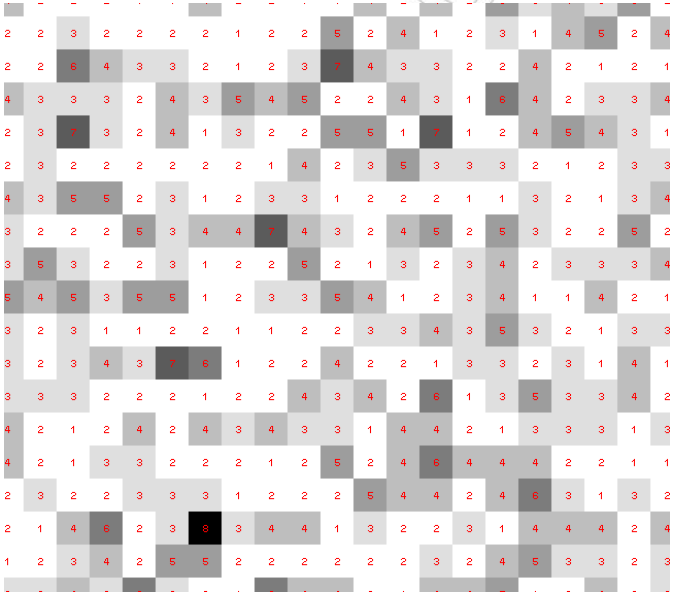
Attenuate beam 4-fold +	360° @ 40 img/s (0.1°/img)	90 s

Are these spots?





Fine φ-slicing + low-intensity data collection make for very weak spots

There is always anomalous signal

Current paradigm

- *Molecular replacement*
- *In case of failure*
  - *Se-Met substitution*
  - *Heavy atom soaking*

Always measure anomalous differences

- *Experimental phasing*
- *Improved MR phases*
- *Identify metal ions*
- *Without extra work*

Always measure 360° of data.

Get best data by

- *Using PILATUS and EIGER detectors*
- *Using fine φ-slicing*
- *Minimizing absolute background*
- *Collecting 360° of low-intensity data*

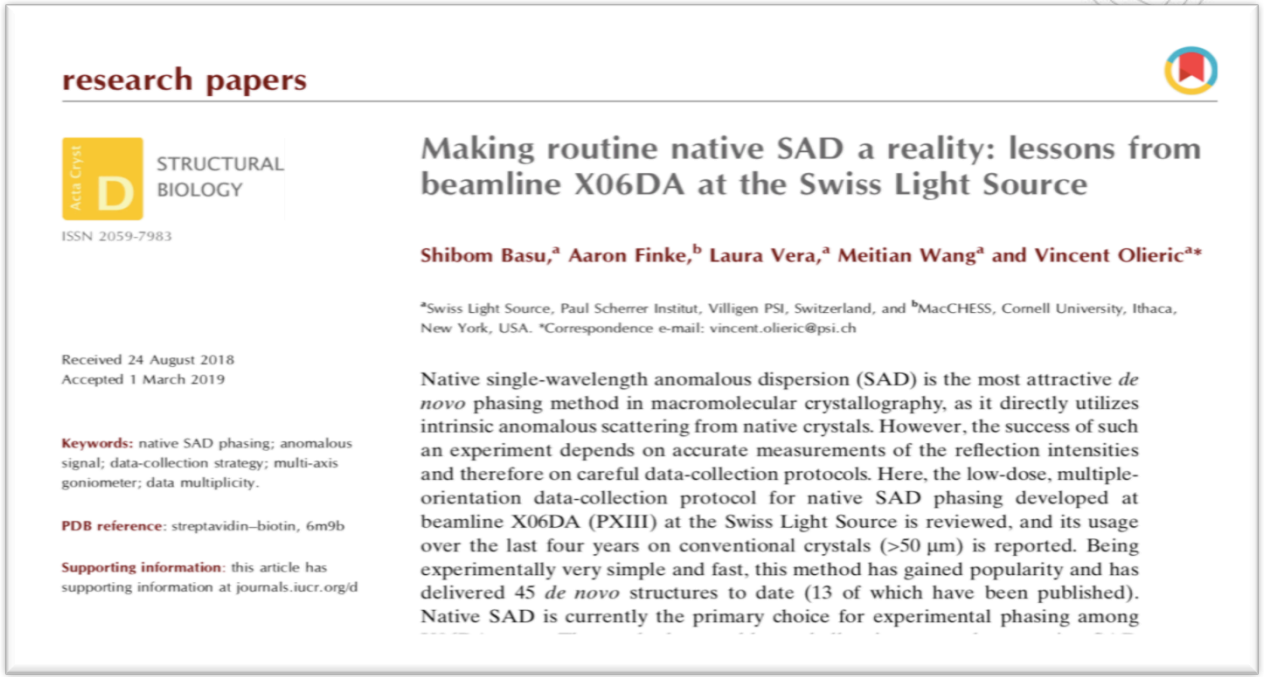
Sophisticated strategies

Collect 360° of data

- *From starting angle suggested by strategy software*
- *With detector at correct distance*
- *With optimized exposure time/attenuation*
- *At non-standard energy (low or high!)*
- *With inverse beam method*
- *From aligned crystal*
- *Multiple times (dose fractionation)*

Never measure less than 360° of data.

Simple native SAD strategy



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STRUCTURAL BIOLOGY

ISSN 2059-7983

Making routine native SAD a reality: lessons from beamline X06DA at the Swiss Light Source

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Received 24 August 2018
Accepted 1 March 2019

Keywords: native SAD phasing; anomalous signal; data-collection strategy; multi-axis goniometer; data multiplicity.

PDB reference: streptavidin–biotin, 6m9b

Supporting information: this article has supporting information at journals.iucr.org/d

Native single-wavelength anomalous dispersion (SAD) is the most attractive *de novo* phasing method in macromolecular crystallography, as it directly utilizes intrinsic anomalous scattering from native crystals. However, the success of such an experiment depends on accurate measurements of the reflection intensities and therefore on careful data-collection protocols. Here, the low-dose, multiple-orientation data-collection protocol for native SAD phasing developed at beamline X06DA (PXIII) at the Swiss Light Source is reviewed, and its usage over the last four years on conventional crystals (>50 µm) is reported. Being experimentally very simple and fast, this method has gained popularity and has delivered 45 *de novo* structures to date (13 of which have been published). Native SAD is currently the primary choice for experimental phasing among

doi.org/10.1107/S2059798319003103

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Maximize anomalous signal

- ***Multiplicity amplifies anomalous signal***
  - Change crystal orientation for true multiplicity
- ***Radiation damage must be avoided***
  - Low-intensity data collection
  - Collect more images at lower dose
- ***Combine data from multiple crystals***

Detector must be free of readout noise.

Get best data by

- *Using PILATUS and EIGER detectors*
- *Using fine φ-slicing*
- *Minimizing absolute background*
- *Collecting at least 360° of low-intensity data*
- *Thinking about your experiment*

Use beam time efficiently

- *Exchange crystal – 20 sec*
- *Center crystal – 20 sec*
- *Collect a dataset – 20 sec*
- *Five-hour shift – 300 min*

Use beam time efficiently

- *Exchange crystal – 20 sec*
- *Center crystal – 20 sec*
- *Collect a dataset – 20 sec*
- *Five-hour shift – 300 min*

- ***Person A :***
  - Collects 82 datasets
  *- **Takes 1 TB of data home***

Use beam time effectively

- *Exchange crystal – 20 sec*
- *Center crystal – 20 sec*
- *Collect a dataset – 20 sec*
- *Five-hour shift – 300 min*

- ***Person A :***
  - Collects 82 datasets
  *- **Takes 1 TB of data home***
- ***Person B :***
  - Thinks about experiment
  - Talks to beamline scientist
  - Has a tea to focus
  - Collects a dataset
  - Has a chat with her mate
  - Collects two more datasets
  *- **Solves structure***

Conclusions

- *Optimize your sample!*
- *With PILATUS or EIGER, collect n × 360° of data with weak beam*
- *With PILATUS or EIGER, Δϕ ≤ ½ mosaicity (XDS) for best data*
- *There is no native data*
- *Ask your beamline scientist / think !*

Read more

- *Förster et al.* ***Transforming X-ray detection with hybrid photon counting detectors***. *Philos Trans A. 2019;377:20180241*.
- *Pflugrath.* ***The finer things in X-ray diffraction data collection.*** *Acta D. 1999;55:1718*.
- *Mueller et al.* ***Optimal fine φ-slicing for single-photon-counting pixel detectors***. *Acta D. 2012;68:42*.
- *Casanas et al.* ***EIGER detector: application in macromolecular crystallography.*** *Acta D. 2016;72:1036.*
- *Winter et al.* ***How best to use photons***. *Acta D. 2019;75:242*.
- *Basu et al.* ***Making routine native SAD a reality: lessons from beamline X06DA at the Swiss Light Source***. *Acta D. 2019;75:262*.
- *Storm et al.* ***Measuring energy-dependent photoelectron escape in microcrystals***. *IUCr J. 2020; 7:129*
- *Förster and Schulze-Briese.* ***A shared vision for macromolecular crystallography over the next five years.*** *Struct Dynam. 2019; 6:064302*.

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Wishing you best data!

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