Light-source diffraction studies of phase transitions under shock loading



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New capabilities for dynamic compression



Shock waves in crystalline materials



Atomic-length scale processes

- Elastic compression
- Plastic deformation
- Phase changes
- Kinetics and metastability

In situ XRD can be critical to determine lattice-level structural information



X-ray diffraction on the ns time scale of shock loading events

LINAC Coherent Light Source (LCLS):

- ~10¹² photons/pulse
- ~100 femtosecond pulses



Advanced Photon Source (APS):

- ~10⁹ photons/pulse
- ~100 picosecond pulses



Image: ANL

Highlight recent experiments:

- Laser-drive compression carbonates at the LCLS
- Gas–gun compression of ZnO & MgF₂ at the APS



Carbonates in the deep Earth



High-P-T phase stability of carbonates is fundamental for understanding the global carbon cycle and carbon storage in the deep Earth



Rapid loading of minerals during impact events



Interpretation of shock metamorphism Understanding role of impact devolatilization

- Solid-state phase transitions
- Melting
- Dissociation
- Dissolution of solid residual phases

 $CaCO_3 \rightarrow CaO^{solid} + CO_2^{gas}$

Laser-based shock experiments are effective in reproducing shock effects observed in naturally shocked minerals



Pump-probe X-ray diffraction at Matter in Extreme Conditions (MEC) beamline









MEC Experiments





SCIENCE

Samples:

Calcite - CaCO₃

- Limestone
- Calcite single crystals

Magnesite - MgCO₃

- Polycrystal (7% porosity)
- Natural gem

Drive Laser:

- 150 & 300-mm phase plates
- 10-15 ns flattop pulse
- Laser energies 10-70 J

XFEL:

- 9.5 & 14 keV
- 20-mm spot size

Line VISAR:

- Wave profiles collected at free surface or sample-LiF interface
- Pressure & shot timing determined via impedance matching 8

Complex polymorphism in calcium carbonate



Rhombohedral $CaCO_3$ -I transitions to a series of low-symmetry phases involving reorientation and tilting of the CO_3 groups



Early calcite shock-wave experiments

Gas-gun studies identified a phase transition ~20 GPa

Based on thermodynamic considerations, high-pressure phase not consistent with the Calcite I-III phases known at the time

Conflicting results concerning degassing with reports ranging between 1-50 mole% devolatilization





Phase transition at 25 GPa



Fig. 7 Preliminary MEC XRD data (8.5 keV) for limestone shocked to 25 GPa. The 2D detector panel is at the top with azimuthally integrated data binufacted antipient calcite and aragonite XRD patterns are shown along with CaCO₂-III-VII patterns



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Release time series from 40 GPa







Future Work:

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Quantitative analysis of liquid scattering (14 keV)



2.8

2.0

1.8

2.2

2.4

d-spacing (angstrom)

2.6

3.0

X-ray densities agree with gas-gun data





Magnesite - MgCO₃

Static:

 Stable up to 100 GPa, above which it undergoes a phase transition to an orthorhombic structure



- Phase change on the Hugoniot near ~100 GPa
- A volume expansion on release was interpreted in terms of decomposition



Sekine et al., 2006



Magnesite-I stable up to melting at >120 GPa



temperature of 2.5-3K at this pressure



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Magnesite Hugoniot

Pressure-density data derived from XRD

Consistent with past gas-gun results

Deviates from calculated Hugoniot based on static results above 90 GPa





Retention of MgCO₃-I structure on release





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ation from X-ray (AS, PCI, ...)



ry

Calcite:

- Crystallographic phase transformation to CaCO₃-VI
- Melting on Hugoniot above 60 GPa
- Reversion to CaCO₃-I on release with no evidence for devolatilization

Magnesite:

- Stable up to melting on the Hugoniot above 120 GPa
- Retention of MgCO₃-I on release



Shock experiments at XFEL:

- Allow us to resolve & differentiate low-symmetry crystal structures
- Provides means of carrying our detailed investigation of release behavior via pump-probe time series



Phase transition in ZnO





High pressure polymorphism in zinc oxide

Fig: adapted from Wang et al. (2018)



- Phase transition a rocksalt phase at moderate pressure (9-16 GPa)
- Transition is common to many wurtzite and zincblende compounds
- Interest in finding routes to quench the ZnO rocksalt phase to ambient conditions due to its favorable optoelectronic properties
- Ultrafast XRD presents a unique capability to study this transformation in real time



Dynamic Compression Sector (DCS): Impact launcher at APS











Multi-frame XRD at DCS

Take advantage of time structure of the synchrotron to collect a series of XRD frames

24-bunch mode well suited to ~100-ns time scale of gun experiments

At DCS collect four frames during the loading and release process for a given shot



Image: ANL





Pink beam X-ray diffraction







Polycrystalline material 20 GPa







Reverberating shock 75 μm ZnO 250 Absorption of ZnO: Thin samples ~75 μ m 200 Transit time ~20 ns 1.5 mm LiF time (ns) LiF impactor 150-Caveat: Can't control x-ray window probe time relative to impact to 100 within 153 ns \rightarrow can't ensure single shock state 50 0 1.2 0.8 1.6

CARNEGIE Science x (mm)

2.0

Single crystal shots in reflection geometry

- Significantly absorbing samples require reflection geometry
- Front surface impact shots: ZnO mounted in Lexan projectile and used as impactor
- Impact TPX/Lexan window
- To optimize 2-theta coverage and reduce low-angle cut off from sample absorption beam comes in at grazing angle 7°









Oriented single crystals 20 GPa



- Large crystallites preserved through the transformation
- Transformation to rocksalt phase with high degree texture



Reproducible transformation textures



Reproducible pattern in terms of where we see the textured diffraction spots

Suggests we capture a reproducible transformation within the time scale of our measurements

ZnO (001)→ TPX

ZnO (001)→ Lexan



Texture analysis \rightarrow orientation relations



Top views of wurtzite phase (a) and rocksalt phase (b) crystal structures.



C-axes loading:



A-axes loading:



RS (200) ~ wurtzite c-axes RS (220) ~ wurtzite a-axes

ZnO conclusions

In situ x-ray diffraction allows for crystallographic verification of the phase transition from $WZ \rightarrow RS$ under shock compression

Single crystals show reproducible transformation textures with strong preferred orientation in transformed rocksalt phase





Phase transitions in MgF₂







Static: rutile \rightarrow CaCl₂ \rightarrow PdF₂



Haines, 2001

Phase transitions in MgF₂ under shock loading



Gun shock data collected at Kumamoto University:

Questions:

What is structure of high-pressure phase?

What structure does the high-pressure phase revert to on release?



36 keV – Single Multilayer Monochromator

Isolate 5th harmonic of U27 (36 keV peak intensity)



- High resolution diffraction peaks in single pulse
- Increased 2θ coverage to determine crystal structure
- High data quality for thick samples & high-Z materials



hemical mechanisms that command the structure-property elation at DCS

However, to our knowledge, the available studies about the Transmission geometry of ZnO are still inconclusive and ontromatical polyer function of ZnO are still inconclusive and ontromatical polyer function of the bulk nodulus (K_0) of the two phases as shown in Table 1, ranging of 135-mm1thick MgE₂ samples with 0.7-mmLiE window 177 to 229 GPa for the B4 phase and from 177 to 229 GPa for the transition pressures of t^{1-1} for m^3 . In ddition, while the transition pressures of t^{1-1} for m^3 and t^{1-1} for m^3 .

ackward phase transitions for nanocrysta cudied,¹⁶ little is currently known about EOS).²² Moreover, the VB4-to-B1 phase ne first principles calculations, the wu ontinuously transform to the rock salt station nexagonal" (*i*-H) pat M_{gF_2} he "tetragonal" neretical computation, Saitta et al.

What is the high pre under shoc

A new capability for x-ray loading at the Dynami

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40 GPa – Modified fluorite phase



Transition to 6+2 coordinated modified fluorite phase







Release and reversion to α -PbO2 + rutile









rutile

 α -PbO₂

MgF_2 Conclusions

- In situ XRD allows for crystallographic verification of the phase transition to modified fluorite phase under plate-impact shock loading
- Reversion to mixture of α -PbO₂ and rutile phase on release
- Demonstration of new capabilities for 36 keV using SMM







GPa/µs



GPa/ns

oading



/100µs

A unique opportunity to investigate a phenomena at various time scales with the same microscopic information from X-ray measurements (XRD, XAS, PCI, ...)

aterials





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SLAC NATIONAL ACCELERATOR LABORATORY

THE Dynamic Compression Sector AT THE ADVANCED PHOTON SOURCE