Wavelength dependence of the damage threshold of inorganic materials under extreme-ultraviolet free-electron-laser irradiation

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We exposed bulk SiC and films of SiC and B₄C to single 25 fs long free-electron-laser pulses with wavelengths between 13.5 and 32 nm. The materials are candidates for x-ray free-electron laser optics. We found that the threshold for surface-damage of the *bulk* SiC samples exceeds the fluence required for thermal melting at all wavelengths. The damage threshold of the *film* sample shows a strong wavelength dependence. For wavelengths of 13.5 and 21.7 nm, the damage threshold is equal to or exceeds the melting threshold, whereas at 32 nm the damage threshold falls below the melting threshold. © 2009 American Institute of Physics. [doi:10.1063/1.3216845]

Free-electron lasers (FELs) produce tunable, coherent, and high-power radiation and have recently been extended to short wavelengths. The FEL in Hamburg (FLASH) produces 25 fs long pulses of more than 10¹² photons in the extreme-ultraviolet (EUV) range between 60 and 6 nm.¹ The Linac Coherent Light Source (LCLS) FEL is scheduled to deliver first light this year.² It will produce 200 fs x-ray pulses with wavelengths between 1.5 and 0.15 nm. The LCLS will be followed by facilities in Germany³ and Japan.⁴

X-ray FELs will enable unique ultrafast scientific research. At the same time, their unique output characteristics put severe requirements on the optics used to guide and shape the x-ray pulses, and the detectors used to characterize them. To avoid damage, various x-ray FEL-material interaction models, damage mechanisms, and optics design guidelines have been suggested.^{5–10} Surface melting is an important physical mechanism since it usually leads to and often sets the threshold for damage. Materials with low atomic number, *Z*, and high melting points are expected to exhibit the highest damage resistance since they absorb less light per unit mass and are mechanically stable. A definitive experimental verification of these models has not been performed since appropriate light sources are not yet available. Instead, existing light sources have been used to test the models by emulating the effect of x-ray FEL radiation,^{11,12} but they are operated typically at wavelengths longer than 32 nm, making extrapolation to x-ray FEL conditions questionable. In this paper we determine the robustness of optical materials that will be used at LCLS, at wavelengths as short as 13.5 nm at FLASH. This wavelength range is substantially closer to the x-ray FEL range than in previous work. Also, we compared bulk materials with thin films and found substantial differences in the damage resistance, suggesting that mechanical

properties and stoichiometry determined by fabrication methodology have a strong effect on the damage resistance.

We studied different low-*Z* materials that are candidates for x-ray FEL optics: (i) SiC slabs fabricated by chemical-vapor deposition with an average grain size of 7.5 μm, (ii) 1 μm thick amorphous SiC films, and (iii) 1 μm thick amorphous B₄C films.¹³ The films were deposited with planar dc-magnetron sputtering systems.¹⁴ The deposition system used for the B₄C films is also being used for the deposition of the coatings on the LCLS x-ray mirrors.¹⁵ In cases (ii) and (iii) the substrates were (100) Si wafers. The B₄C films deposited under regular conditions had high compressive stress (>2 GPa), which led to delamination from the substrate. We increased the B₄C deposition pressure to reduce the stress (to 0.9 GPa), at the expense of a slight increase in high-spatial-frequency roughness (HSFR). Grains appear to be separated by microvoids in some areas. Atomic force microscopy (AFM) measurements on other B₄C films deposited under identical conditions, but with different thicknesses indicated that the grain size and resulting HSFR scale with film thickness. This type of morphology is consistent with a “zone 1/zone T” structure model proposed by Thornton¹⁶ and is also consistent with data on sputtered B₄C films presented in earlier literature.^{16–18} X-ray photoelectron spectroscopy on a B₄C film aged for about 1 month indicated that the top 9 nm of the film is oxygen and carbon rich (64% B, 22% C, and 13% O, atomic) with the oxygen and carbon concentrations rapidly diminishing with depth from the top surface. Rutherford backscattering (RBS) measurements across the entire B₄C film thickness indicated atomic ratio of boron-to-carbon=3.7 (near perfectly stoichiometric), with 6% oxygen present, believed to come from the B₄C sputtering target. Using the RBS measurements combined with scanning electron microscopy (SEM) thickness and EUV reflectance measurements, it was determined that the density of

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FIG. 1. Representative SEM images of bulk SiC exposed to the FLASH beam slightly above the damage threshold at wavelengths of 13.5, 21.7, and 32 nm, respectively. The widths of the images are (a) 10.6 μm , (b) 25.1 μm , and (c) 24.5 μm .

the films is 2.28 g/cm^3 (about 90% of crystalline B_4C). The aforementioned properties are generally consistent with the properties of B_4C films made using physical vapor deposition methods. RBS measurements on SiC films deposited under conditions similar to the 1 μm thick SiC films used here indicated an atomic ratio of silicon-to-carbon of 0.98 (near-perfectly stoichiometric), with 5% oxygen. The density of the SiC films was determined to be 2.98 g/cm^3 (about 93% of crystalline SiC).

We exposed the different samples to single pulses of the focused FLASH beam under normal incidence. FLASH was operated at wavelengths of 32, 21.7, and 13.5 nm. The pulse energies were up to $\sim 10 \mu\text{J}$, and the pulse duration was 25 ± 5 fs. The FLASH beam was focused onto the sample using a grazing incidence ellipsoidal mirror to a Gaussian beam radius of 5 to 10 μm . The pulse energy was adjusted using a gas attenuator, and a gas monitor detector (GMD) was used to measure the energy of each pulse.¹⁹ In order to extract damage thresholds, we found it necessary to study the onset of surface modification at low pulse energies at or below 1 μJ . Since the readout electronics of the GMD was not optimized for such low pulse energies we were only able to record the energy averaged over multiple pulses. Therefore, we used a statistical method to obtain the damage thresholds as for previous experiments.¹¹ The statistical pulse energy distribution of a self-amplified-spontaneous-emission FEL follows a Gamma distribution with a shape parameter M that describes the number of optical modes in the radiation source.²⁰ Fitting a Gamma distribution to the measured high-energy (weakly attenuated) pulses, we found values for the shape parameter M of 1.5, 1.5, and 3.5 for wavelengths of 13.5, 21.7, and 32 nm, respectively. We exposed multiple spots (15–39) to pulses at a certain gas attenuator setting, corresponding to a certain average pulse energy. Using AFM, SEM, and white-light interference microscopy,²¹ we determined the fraction of the spots that are visibly undamaged. Finally, we determined the damage threshold fluence interval by mapping the undamaged and damaged spots onto the statistical pulse energy distribution.

We determined the Gaussian beam radii from the scaling of the measured damaged area as a function of the incoming pulse energy.^{22,23} We found radii of 4.8, 8.1, and 8.3 μm at wavelengths of 13.5, 21.7, and 32 nm, respectively.

We used white light interference microscopy²¹ to determine surface height profiles. Figure 1 shows representative profiles for bulk SiC exposed to the FLASH beam slightly above the damage threshold at wavelengths of 13.5, 21.7, and 32 nm, respectively. This shows that the focal spot of the beam is well defined, allowing an accurate determination of the fluence. The grains visible in the images are of order the spot size. The three images look similar, indicating that the

TABLE I. Summary of the exposure runs and the extracted damage thresholds.

Material	Wavelength (nm)	Number exposures/damage sites	Lower/upper fluence (mJ/cm^2)	Damage threshold (mJ/cm^2)
SiC film	13.5	35/2	206/282	326
		35/6	380/434	
	21.7	39/14	161/172	178
		15/4	186/228	
B_4C film	32	39/6	33/36	35
		35/12	323/350	
	13.5	35/5	333/388	348
		39/17	96/102	
		39/10	72/79	
		15/3	142/181	
SiC bulk	32	39/1	18/23	18
		39/2	22/26	
	13.5	39/11	10/10	685
		35/21	663/707	
		39/21	321/339	
		39/34	254/277	
	21.7	39/22	75/77	141
		20/8	54/58	

damage mechanism is likely to be the same. For the SiC film and the slab samples, we detected only craters, whereas for the B_4C films we observed both extrusions, primarily at low pulse energies, and craters, primarily at large pulse energies. At a wavelength of 32 nm, we detected only extrusions on B_4C films for pulse fluences of up to 187 mJ/cm^2 . At a wavelength of 21.7 nm, we detected primarily extrusions up to 728 mJ/cm^2 , and a few craters between 243 and 1213 mJ/cm^2 in the B_4C films. Finally, at a wavelength of 13.5 nm, we detected only extrusions in the B_4C films up to 963 mJ/cm^2 . The extrusions might be caused by a phase change in analogy to similar volume expansions studied in amorphous-carbon films.²⁴

We consider the surface damaged if we detect any surface modification by SEM, AFM, or white light interferometry. Table I lists the exposure runs that we performed along with the fraction of exposed sites identified as damaged. From this data, we can extract a range for the damage threshold using the statistical model for each exposure run, also listed in Table I. Finally, we take the damage threshold as the average of the midpoints of these ranges, weighted by the number of exposures per run.

We use the local dose as the basis for extrapolating FLASH results to the hard x-ray regime. This concept is supported by the fact that in both cases the absorption mechanism is photoionization. The electrons then cascade down to energies of order 10 eV before interacting with the lattice ions. In both cases, the timescale for heat conduction is much longer than the radiation pulse lengths, supporting the assumption that the absorbed energy stays confined and ends up as thermal energy before damage processes begin. The calculation of the dose from the incident fluence accounts for the particular absorption mechanisms that act in the two different wavelength regimes (i.e., K -shell, L -shell, etc.). For the FLASH case, the threshold surface dose is obtained by dividing the damage threshold fluence by the attenuation length. We assume that the optical properties of the materials do not change during the pulse²⁵ and that the materials are homogeneous, in which case the maximum dose occurs at the surface. Table II shows the attenuation

TABLE II. Summary of damage threshold doses.

Wavelength (nm)		13.5	21.7	32	Theor.
Attenuation length (nm)	Bulk SiC	224	97	48	
	SiC film	241	105	52	
	B ₄ C film ^a	232	91	45	
	B ₄ C film ^a	130	54	28	
Damage threshold (eV/atom)	Bulk SiC	2.0	2.0	1.9	1.00
	SiC film	0.9	1.2	0.5	1.00
	B ₄ C film	0.8	0.5	0.2	0.77
	B ₄ C film	1.4	0.8	0.3	0.77

^aWe used the actual measured stoichiometry.

lengths and damage threshold doses for bulk SiC, SiC films, and B₄C films as a function of wavelength. We estimate the error in the damage threshold to be about 50% due to errors in the beam area, the energy measurement of the gas detector, and the small number of exposures per exposure series. Thermodynamic calculations show that crystalline SiC reaches the melting temperature at a dose of 0.63 eV/atom.²⁶ The heat of fusion is 0.37 eV/atom,²⁶ so that a dose of 1.0 eV/atom is required to melt the material. Since the measured damage threshold is 2 eV/atom, bulk SiC exhibits a remarkable resistance to single-pulse damage in the EUV regime. The rather weak wavelength dependence of the damage threshold dose over the range of 13.5–32 nm gives hope that the damage threshold at 1.5 nm, (the long wavelength limit of LCLS) is similar.

Both the SiC and the B₄C films show a strong wavelength dependence of the damage threshold dose. The threshold at 32 nm is particularly low, even when we correct for the actual stoichiometry of the B₄C films. It is possible that the observed low damage thresholds are a manifestation of uncertainties in the absorption coefficients of sputtered SiC and B₄C films containing oxygen, especially in the long wavelength regime. Crystalline B₄C has a melting dose of 0.66 eV/atom. Both the SiC and B₄C film damage thresholds are at or above the melting thresholds of the corresponding crystalline materials at 13.5 and 21.7 nm wavelengths, but not as high as the bulk SiC samples. Since many grazing incidence optics for XFELs will use thin coatings, the lower damage thresholds doses found for thin films should be used as guidance for these facilities.

In summary, we measured the single-pulse damage resistance of candidate materials for LCLS optics at the FLASH facility, including bulk SiC and SiC and B₄C films. We found that the damage thresholds at 13.5 and 21.7 nm wavelengths are above the melting thresholds, giving hope that a similar threshold or better can be expected at initial LCLS operation at a wavelength of 1.5 nm. At 32 nm, the damage thresholds of the films are below the melting threshold, which may be attributed to increased absorption.

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