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EELS characterization of zirconium hydrides

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Abstract. — The energy of the first plasmon loss peak has been determined for the ϵ -Zr hydride phase in pure Zr. The peak energy is sufficiently different from that of the δ -hydride to permit EELS “fingerprinting” with ease using a parallel EELS system, but with greater difficulty using serial EELS, because of drift associated with the slower collection rates. Experiments on Zr-2.5Nb alloys showed that the hydride phases can also be distinguished using EELS, facilitating the analysis of hydride precipitates and blisters in nuclear reactor pressure tubes.

1. Introduction.

A major application of the electron energy loss spectroscopy (EELS) technique is to analyze for elements of low atomic number that are not detectable by conventional thick-window energy dispersive X-ray spectrometers. Analysis is most commonly performed using the characteristic edges caused by inner-shell ionizations. However, the low-loss peaks that are generated by plasmons (collective oscillations of the valence electrons) have been shown to be useful for quantitative microanalysis in Al alloys [1] and for phase identification [2].

Zaluzec *et al.* [3, 4] have shown that Zr, Ti and Mg hydrides give rise to plasmon peaks having characteristic energies that are larger than those of the parent metal. The energy shift may be explained in terms of a change in the density and effective mass of the “free” valence electrons associated with hydride formation.

Because there are three distinct hydrides in both Zr and Ti, a study was initiated to investigate the possibility of identifying the various hydride species using plasmon EEL spectra. Resolvable shifts in plasmon loss energies of γ - and δ -hydrides relative to the parent metal (Ti and Zr) were observed, and used for “fingerprinting” the phases [5, 6]. These results also showed that the Zr and Ti hydride plasmon energies reported in references [3] and [4] were originated from Zr γ -hydride and Ti δ -hydride respectively. In this article, we describe an extension of the EELS plasmon loss

energy technique to the ϵ -hydride in the Zr-H system and demonstrate a practical application to the characterization of the major phases in hydrided Zr-2.5Nb. Practical applications would include (a) the rapid identification of hydrides that are responsible for delayed hydride cracking by precipitation of crack tips, and (b) the identification of hydride phases in blisters formed in Zr-2.5Nb. Hydride blisters are accumulations of hydride phase that may form at the cold end of a temperature gradient [7]. They were observed in Zircaloy-2 (Zr-1.15 at% Sn-0.25 at% Fe-0.18 at% Cr-0.08 at% Ni) pressure tubes in CANDU (Canadian Deuterium Uranium) nuclear reactors that had been in contact with its calandria tubes [8].

Zr hydride can form in one of three crystal structures, depending on hydrogen concentration and cooling rate from solution temperature [9-15]. At low to medium concentrations, face-centred-tetragonal (fct) γ -ZrH with $c/a > 1$, and face-centered-cubic (fcc) δ ($\sim \text{ZrH}_{1.5}$) are observed. γ -hydride is metastable and its formation is favoured by fast cooling rates [15], whereas δ -hydride is the equilibrium phase. ϵ -hydride forms at a hydrogen concentration of greater than ≈ 62.5 at%, and has a fct crystal structure but with $c/a < 1$.

2. Experimental details.

Binary zirconium alloys containing 0.36, 44.3, 62.8, 64.3, and 64.8 at% H were prepared with the aid of pressure-temperature isotherms, as described in reference [16]. Zr-2.5Nb material was charged with 44.0 at% H at 925 K from which the reaction was stopped by quenching the specimen tube in a Dewar flask of liquid nitrogen. Another Zr-2.5Nb specimen was charged with 65.5 at% H (specimen G169) but in a different manner; pure hydrogen was created by heating uranium hydride slowly, so the Zr-2.5Nb specimen absorbed an increasing amount of hydrogen while being heated to 1123 K. The specimen was then slowly cooled to room temperature, while, at the same time, the hydrogen pressure was decreased through a reduction in the temperature of the uranium hydride turnings. Thin foils of Zr-H and Zr-2.5Nb-44.3 at% H specimens were made using a Metal Thin electropolishing machine. Foils of Zr-2.5Nb containing high hydrogen concentrations were brittle and difficult to electropolish to electron transparency. They generally thinned uniformly instead of being electropolished with a dimpled profile. A modified technique was employed. A relatively thick disc (≈ 0.3 mm) was carefully dimpled on both sides with a mechanical dimple grinder. Approximately 0.07 mm was left in the middle of the disc which was then electropolished with an automatic Metal Thin electropolishing machine. EEL spectra were obtained using either a Gatan serial EEL spectrometer (model 607) or a Gatan parallel EEL spectrometer (PEELS model 666). The resolution of the EEL spectrometer, defined by the full-width half-maximum of the zero loss peak, varied from 0.9 eV with an unsaturated LaB₆ filament operating at an accelerating voltage of 120 keV to 2.8 eV when operating with a saturated LaB₆ at 300 keV. The highest resolution was employed for EEL spectra with ϵ -hydride because its plasmon loss energy was observed to be only two or three tenths of an eV from that of δ -hydride. All the low loss EEL spectra were calibrated using the plasmon peak of pure Al, which was taken to be 15.3 eV [1].

3. Plasmon loss energies.

Electron-transparent foils from Zr-0.36 at% H and Zr-44.3 at% H were employed to obtain the low loss regions of the EEL spectra for Zr, γ - and δ -hydrides using the serial EEL spectrometer. The 0.36 at% H specimen, identical to that used by Weatherly [17], contained γ -hydrides in a Zr matrix. The 44.3 at% H specimen contained all three phases, Zr, γ - and δ -hydrides. This

observation is in agreement with the general accepted opinion that γ -hydride is a metastable phase produced by a martensitic shear transformation. Plasmon loss energies determined from EEL spectra of these specimens have been reported previously [4, 5], and are summarised in table I. The 62.8 at% H specimen contained γ - and δ -hydrides; two values each were obtained for the plasmon loss energies for each phase. These were 18.5 and 19.3 eV for γ - and δ -hydrides respectively.

Table I. — *Experimental plasmon loss energies in Zr-H specimens.*

Specimen	Plasmon Loss energy (in eV)			
	Zr	γ -hydride	δ -hydride	ϵ -hydride
0.36 at% H	$(16.8 \pm 0.2)_{10}$	$(18.3 \pm 0.2)_{16}$	$(19.2 \pm 0.2)_{14}$	serial EELS
44.3 at% H				serial EELS
62.8 at% H		$(18.5)_2$	$(19.3)_2$	serial EELS
64.3 at% H	(Serial EELS)			$(19.6 \pm 0.5)_{10}$
	(PEELS)			$(19.6 \pm 0.1)_8$
64.8 at% H	(PEELS)			$(19.6 \pm 0.1)_{15}$

Error limits represent 2 standard deviations; number of experimental measurements is given as subscript

Fig. 1. — EEL plasmon spectrum of ϵ -hydride in Zr-64.3 at% H; plasmon loss peak at 19.6 eV.

Initial attempts to distinguish the plasmon loss energy of ϵ -hydride relative to δ -hydride using serial EELS yielded inconclusive results. Further work showed that the distinction could be made if suitable collection conditions were employed. To minimize drift of the electron beam energy, it was necessary to use a very short dwell time, 10 msec/channel, with the intensity of the zero-loss peak low enough to avoid saturating the detector. Ten spectra from ϵ -hydride of the 64.3 at% H alloy were collected and analyzed; the plasmon loss peak was at (19.6 ± 0.5) eV, with the error limits representing two standard deviations.

In contrast to serial EELS, unambiguous measurements were made readily using the high collection efficiency of the PEELS system. A total of twenty two spectra for ϵ -hydride were collected (using dwell times of 100 to 500 ms) for the 64.3 at% H and the 64.8 at% H specimens. One such EEL spectrum is shown in figure 1. The plasma loss peak was at (19.6 ± 0.1) eV in both cases. By comparing the plasmon loss energies for ϵ -hydride as determined by serial and parallel EELS (Tab. II), it is clear that PEELS offer better accuracy and reproducibility. The error limits from the serial EELS measurements also make it impossible to distinguish ϵ -hydride from δ -hydride unless a large number of measurements are made.

Table II. — *Experimental plasmon loss energies of hydrides in Zr-2.5Nb.*

Specimen	Plasmon Loss energy in (eV)		
	γ -hydride	δ -hydride	ϵ -hydride
44.3 at% H	$(18.3 \pm 0.2)_8$	$(19.2 \pm 0.2)_6$	
65.5 at% H			$(19.6 \pm 0.1)_{10}$

The microstructure of the Zr-2.5Nb-44.0 at% H specimen consists of martensitic grains (Fig. 2a) and clear grains with or without twinned structures (e.g. Fig. 2b). The grain diameters were 2 to 10 μm . Analysis of electron microdiffraction patterns from the clear grains revealed that they were fcc γ -hydride, with a lattice parameter of 0.478 nm. The plasmon loss energy, as determined from six EEL spectra, was (19.2 ± 0.2) eV. The twinned structures were shown by electron microdiffraction to be fct γ -hydride, with $a = 0.464$ nm, and $c = 0.498$ nm. The plasmon loss energy was (18.3 ± 0.2) eV, analyzed from eight EEL spectra. In figure 3 are examples of EEL spectra for γ - and δ -hydrides in Zr-2.5Nb respectively. Their plasmon energies are indistinguishable from those for Zr-H alloys, despite the alloying addition of Nb.

TEM/AEM examination of the Zr-2.5Nb-65.5 at% H specimen (G169) revealed " β -Zr"-like filaments in a matrix of solid ϵ -hydride (Fig. 4). These filaments had widths of about 20 nm, and were spaced 0.1 to 0.3 μm apart. Energy dispersive X-ray analysis revealed a Nb concentration of greater than 15 at% in the phase. Attempts to obtain good microdiffraction patterns from them were unsuccessful. Because hydrogen addition was carried out at 1123 K in the $(\alpha + \beta)$ -field while the specimen absorbed an increasing amount of hydrogen during heating, the filaments were thought to be hydrogen-stabilized β -Zr phase. The matrix contained about 0.7 at% Nb. Electron microdiffusion analyses were consistent with the matrix being fct ϵ -hydride phase, with $a = 0.492$ nm, and $c = 0.456$ nm. Ten EEL spectra (e.g. Fig. 5) were collected with the PEELS system, and they were analyzed to give a plasmon loss energy of (19.6 ± 0.1) eV. This energy is

Fig. 2. — Microstructure of Zr-2.5Nb-44.0 at% H, showing (a) martensitic grain structure, and (b) δ -hydride grain with twinned γ -hydride.

Fig. 3. — EEL plasmon spectra of γ - and δ -hydrides in Zr-2.5Nb-44.0 at% H showing plasmon loss energies of γ -hydride at 18.4 eV and δ -hydride at 19.3 eV.

consistent with that for ϵ -hydride in Zr-H specimens. A summary of the plasmon energies for γ -, δ - and ϵ -hydrides is given in table II.

Fig. 4. — Microstructure of Zr-2.5Nb-65.5 at% H showing “ β -Zr” like filaments in an ϵ -hydride matrix.

Fig. 5. — EEL plasmon spectrum of ϵ -hydride in Zr-2.5Nb-65.5 at% H; plasmon loss peak at 19.6 eV.

The lattice parameters of all the hydrides in Zr-H and in Zr-2.5Nb-H alloys, measured using electron microdiffraction patterns, are summarized in table III. Literature values are also listed for comparison.

Table III. — *Lattice parameters of hydrides in Zr-H and Zr-2.5Nb-H alloys.*

Specimen	γ -hydride	δ -hydride	ϵ -hydride
Zr-H alloys			
0.36 at% H	a = 0.4596 nm c = 0.4969 nm (from [16])		
44.3 at% H	a = 0.460 nm c = 0.497 nm	a = 0.478 nm	
Literature	a = 0.458 to 0.460 nm; c = 0.497 nm [10-12]	a = 0.478 to 0.482 nm [9,11,12]	
64.3 at% H			a = 0.490 nm c = 0.460 nm
64.8 at% H			a = 0.492 nm c = 0.461 nm
Literature			a = 0.476 to 0.494 nm; c = 0.472 to 0.446 nm [9,11,12]
Zr-2.5Nb-H alloys			
44.0 at% H	a = 0.464 nm c = 0.498 nm	a = 0.478 nm	
65.5 at% H			a = 0.492 nm c = 0.456 nm

4. EELS 'fingerprinting'

The calibrated plasmon loss energies listed in tables II and III can be used for 'fingerprinting' hydride phases in Zr and Zr-Nb. This technique has been shown to be successful in phase identification in a specimen of Zr-44.3 at% H where all three phases, Zr, γ -hydride co-exist (Fig. 6). EEL spectra were taken from each of the regions to locate the plasmon loss peaks. Based on these peak values, the phases corresponding to each region were identified and subsequently verified with microdiffraction results [4]. It would have been possible to identify the hydride species by

electron microdiffraction, but the EELS fingerprinting technique is very much faster, because an EEL spectrum can be acquired and analyzed on-line in minutes, compared to several hours by electron microdiffraction.

Fig. 6. — Regions analyzed in a thin foil of Zr-44.3 at% H and identified after the EELS results.

The EELS fingerprinting technique is best illustrated for hydride phase identification in hydride blisters grown on Zr-2.5Nb. The information can be used as a basis for comparison with predictions from the blister growth model. Typically the centre of a blister older than 3 months is made up of δ -hydride, while the phases at the edge are mixtures of γ - and δ -hydrides. An example of a fine-grained γ -hydride region at the edge of a blister on Zr-2.5Nb pressure tube aged for seven months is shown in figure 7. Because of the complexity of the microstructures of blisters, which contain high dislocation densities and sometimes twins, the EELS fingerprinting technique is clearly superior to electron diffraction for routine analyses. Moreover, EELS has the advantage of being able to probe into very fine grains, $\sim 0.5 \mu\text{m}$, in Zr-2.5Nb pressure tube, while it would be very difficult to obtain microdiffraction patterns from such small areas.

5. Conclusions.

This article describes an EELS plasmon fingerprinting technique used in identifying Zr, γ - and δ -hydrides and extended to ϵ -hydride. The plasmon loss energy of ϵ -hydride was found to differ by only about two-tenths of an eV from that of δ -hydride. Experiments revealed that parallel EELS offered better accuracy and reproducibility than serial EELS in detecting such small energy difference. The energy peaks will be useful for fingerprinting phases in solid hydride specimens and in hydride blisters grown on Zr-2.5Nb pressure tube.

Fig. 7. — File-grained γ -hydride region in an aged blister on Zr-2.5Nb.

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