DLI-MOCVD Crx Cy coating to prevent Zr-based cladding from inner oxidation and secondary hydriding upon LOCA conditions

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J.C. Brachet, S. Urvoy, E. Rouesne, G. Nony, M. Dumerval, et al.. DLI-MOCVD Crx Cy coating to prevent Zr-based cladding from inner oxidation and secondary hydriding upon LOCA conditions. Journal of Nuclear Materials, 2021, 550, pp.152953. 10.1016/j.jnucmat.2021.152953 . hal-03212406

HAL Id: hal-03212406

<https://hal-ensta-bretagne.archives-ouvertes.fr/hal-03212406>

Submitted on 29 Apr 2021

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Title: “DLI-MOCVD Cr_xC_y coating to prevent Zr-based cladding from inner oxidation and secondary hydriding upon LOCA conditions”

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ABSTRACT:

Zirconium-based claddings with an outer chromium coating resistant to corrosion are studied and developed as an evolutionary Enhanced Accident Tolerant Fuel (E-ATF) concept for light water reactors. However, in hypothetical LOss-of-Coolant-Accident (LOCA) conditions, following clad ballooning and burst, the outer coating does not allow to protect the inner surface of the cladding from High Temperature (HT) steam oxidation and associated secondary hydriding due to steam starvation occurring within the gap between the clad inner surface and the nuclear fuel pellets.

To address this issue, DLI-MOCVD (Direct Liquid Injection of Metal-Organic precursors - Chemical Vapor Deposition) Cr_xC_y coatings have been developed and successfully deposited onto the inner surface of Zr-based cladding tube prototypes. Then, preliminary two-sided oxidation tests have shown that such inner coating is able to increase the resistance to oxidation at HT of the inner clad surface.

The present study aimed at performing new steam oxidation tests at 1200°C on Zircaloy-4 clad prototypes with a 5-20µm-thick Cr_xC_y inner coating, in conditions more representative of LOCA, after a first internal pressure-induced burst step. Additionally, complementary two-sided steam oxidation tests have been carried out up to 1h at 1200°C, on short inner and/or outer-coated clad segments. Finally, Post-Quench (PQ) Ring Compression Tests (RCTs), fractographic analysis and deep metallurgical investigations including neutron-tomography have been performed to get more insights into the PQ behavior of the inner-coated clad.

Among other results, it is shown that the inner Cr_xC_y coating makes it possible to reduce significantly the oxidation and the associated secondary hydriding of the clad inner surface, after ballooning and burst. After at least 600s under steam at 1200°C, the reference uncoated clad fails upon final water quenching while the inner-coated prototype keeps its integrity. PQ RCTs showed a higher strength of the inner-coated material, related to lower oxygen and hydrogen uptakes of the substrate.

KEYWORDS: Enhanced Accident Tolerant Fuel, Cr_xC_y DLI-MOCVD inner coating, Zircaloy-4, LOCA, high temperature steam oxidation, secondary hydriding, post-quenching strength and ductility

HIGHLIGHTS:

- 5-20µm-thick Cr_xC_y coating deposited by DLI-MOCVD onto inner surface of Zircaloy-4 cladding,
- Protection against steam oxidation at 1200°C comparable to outer PVD-deposited 10-15µm-thick Cr coating.
- Significant reduction of secondary hydriding after clad ballooning and burst.
- Higher mechanical strength of the cladding, upon and after water quenching.

1. INTRODUCTION

Zirconium-based claddings with outer corrosion resistant chromium-based coatings are studied and developed worldwide as an evolutionary Enhanced Accident Tolerant Fuel (E-ATF) concept for light water reactors, bringing additional High Temperature (HT) oxidation resistance in hypothetical accident conditions [1]-[23]. However, in case of leaking fuel or following clad ballooning and burst upon hypothetical Loss-of-Coolant-Accident (LOCA), the outer coatings developed so far are not able to protect the clad inner surface from steam oxidation and from the associated secondary hydriding phenomena. To address this issue, DLI-MOCVD (Direct Liquid Injection of Metal-Organic precursors - Chemical Vapor Deposition) Cr_xC_y inner coating has been developed and successfully deposited on the inner surface of Zr-based cladding tube prototypes [24]-[26]. Then, preliminary oxidation tests have shown that such inner coating may increase the oxidation resistance of the clad inner surface at HT [27]-[28]. However, the preliminary two-sided HT oxidation tests carried out until now were not fully representative of LOCA situations, in which the clad inner surface is exposed to an oxidizing steam atmosphere confined into the gap between the nuclear-fuel pellets and the cladding tube. In such conditions, due to steam starvation, the cladding materials experience localized massive hydrogen uptake, characterized by “hydrogen peaks”, a few centimeters from the burst opening, with hydrogen contents up to 3000-4000 parts per million in weight (wt.ppm). Such phenomena may induce cladding embrittlement and loss of clad integrity upon the final water quenching and/or in Post-Quenching (PQ) situation. Thus, they have been extensively studied at ANL [29]-[30], at JAERI/JAEA [31]-[39], at KIT [40]-[46] and at other institutes [47]-[48]. Additionally, modelling efforts have been carried out [50]-[52] and safety criteria have been proposed in some countries, to account for this clad embrittlement due to secondary hydriding, following ballooning and burst upon LOCA conditions [53]-[57].

To address this issue, the present study aimed at performing new and more LOCA-representative HT oxidation tests on the DLI-MOCVD inner Cr_xC_y -coated Zr-based cladding prototypes [27]-[28]. Additionally, complementary HT two-sided steam oxidation tests have been carried out on inner and/or outer-coated short clad segments. Finally, PQ Ring Compression Tests (RCTs), fractographic analysis and deep metallurgical investigations (including neutron tomography to obtain 3D mapping of hydrogen) have been performed to get more insights into the PQ behavior of inner-coated clad.

2. MATERIALS AND EXPERIMENTAL PROCEDURES

2.1. Coated and uncoated materials

Uncoated substrate materials:

Fully recrystallized Zircaloy-4 with high tin and oxygen contents (HSn-HO-Zirc-4), 0.6 mm-thick, cladding tube has been used as a substrate. The contents of the main alloying chemical elements are given in Table 1. It has to be mentioned that the quality of the “old grade” Zircaloy-4 cladding material tested is not representative of modern industrial advanced cladding materials in as-received conditions. The clad inner surface exhibits a high roughness and some superficial oxidation, as shown below. Thus, in the following, this material has to be considered as having a “lower-bond quality” compared to modern nuclear fuel cladding materials, especially at its inner surface. Indeed, it is likely that this poor quality (rough and slightly oxidized) of the as-received clad inner surface negatively impact the adhesion of the inner coating.

Alloy	Wt.%				Wt.ppm	
	Sn	Fe	Cr	O	H	N
HSn-HO-Zirc-4	1.62	0.20	0.10	0.18	12	60

Table 1 - Contents of the main alloying chemical element of the “old grade” high tin – high oxygen Zircaloy-4 used as a substrate

Coated materials:

Cr_xC_y coatings were deposited directly inside a bundle of 16 cladding tubes with a horizontal hot-wall CVD chamber of approximately 0.15 m of inner diameter and 1 m of length. A liquid solution of bis(ethylbenzene)chromium (precursor) and toluene (solvent) was vaporized using a Kemstream DLI system and sent to the inside of the 16 claddings segments at 670 Pa heated at 325°C by a tubular furnace. More information about the deposition chamber can be found in [25].

A schematic view of the inner-coated 1m-long Zircaloy-4 cladding tube prototype studied here is shown in Figure 1. In this figure, the different samples machined and tested are represented together with the as-received inner coating thicknesses measured by Optical Microscopy (OM) at different axial locations. Due to the deposition process used, there is an axial gradient of the inner coating thickness from the left end to the right end of the tube. In consequence, depending of the axial position, the inner Cr_xC_y coating thickness ranges from ~20µm at the M2 location down to ~6µm at the M5 location. It is nevertheless possible to inject sequentially the reactive vapor at each end of the chamber to smoothen thickness profiles, as demonstrated in [24] for a single 1m-long cladding segment. In the following, one has to account for the different inner coating thicknesses when analyzing the results. The different clad segment geometries tested and/or analyzed are:

- “Mi” 5mm-long rings, for as-received microstructural observation and hydrogen analysis;
- “Oi” 30mm-long clad segments for two-sided HT steam oxidation testing followed by PQ microstructural and micro-chemical analysis and RCTs. Among these samples and following the inner DLI-MOCVD deposition process, ~10µm-thick outer chromium coating has been deposited on two of these 30mm-long segments by the PVD process already used for previous CEA-Framatome-EDF studies on outer Cr-coated advanced nuclear fuel claddings [2][7][11][20][21]. Following preliminary air oxidation tests carried out at 1100°C on such double-coated claddings [28], the idea here was to have a deeper view of the benefits, regarding oxidation resistance and PQ mechanical behavior, of double (inner + outer) coating protection in conditions more representative of LOCA, *i.e. two-sided HT steam oxidation at 1200°C followed by direct water quenching*.
- “E1” and “E2” longer segments that have been used for two-steps LOCA testing, including a first internal pressure thermal ramp test leading to cladding burst and a subsequent steam oxidation test at 1200°C followed by direct water quenching. So, these two particular segments have been specifically devoted to the study of secondary hydriding following burst occurrence upon LOCA. The results have been compared to those obtained in the same conditions on the uncoated clad.

2.2. Experimental procedures

Internal pressure burst tests, one/two-sided HT steam oxidation tests, PQ mechanical tests and metallurgical examination including micro-chemical analysis and neutron radiography/tomography have been carried out on several CEA’s facilities already described in details in [47] and [58]-[62]. They are only briefly recalled below.

- Internal pressure thermal ramp tests to burst have been carried out using the “EDGAR” facility used at CEA for more than twenty years to perform thermal-mechanical tests upon LOCA conditions on 500mm-long clad segments [58]. Direct Joule heating was used to be able to achieve fast enough heating rates with good circumferential temperature homogeneity. The evolution of the cladding outer diameter was measured using a non-contact laser extensometer. Temperature was monitored and controlled by pyrometers and internal thermocouples. The tests were performed under steam and stopped just after the burst. The tested clad segments were then cooled at moderate rate (*a few tenths of °C/s at the beginning of the cooling*) down to RT. The internal pressure and heating rate applied here were set to 40 bars and 25°C/s respectively. These particular values have been chosen to allow a comparison with previous data obtained on uncoated M5_{Framatome} reference cladding segments [47]. Since the final cladding hoop strain at burst and the overall balloon size, and thus the inner gap size, may depend on the internal pressure and heating rate applied, the results obtained here should not be extrapolated directly to other testing conditions. Indeed, due to the limited ductility of the inner coating, it is likely that larger clad deformations than those experienced here promote cracking of the inner CrC coating, and thus influence secondary hydriding).

- Following the creep and burst sequence, 150mm-long segments have been machined from the segments tested in “EDGAR” by localizing the burst opening at their mid-length. An alumina rod (8.2mm in diameter) has been inserted inside the tube to simulate the gap between the fuel pellets and the clad inner surface. Zircaloy-4 end caps have been welded at both clad segment extremities. A typical macrograph of a pre-burst clad segment before the subsequent HT oxidation test is shown in Figure 2-(a).

- Oxidation tests in steam at 1200°C for different oxidation times have been carried out in the CEA’s “DEZIROX 1” facility [59]-[61]. Upon the HT oxidation process, the axial temperature gradient along the 150mm-long clad segment is lower than 10-15°C. Direct water quenching from 1200°C down to Room Temperature (RT) was applied at the end of the isothermal HT oxidation step. It must be mentioned that, compared to a direct quenching from the HT β_{Zr} temperature range, when the final water quenching is applied at 600 or 700°C (after a first cooling at an intermediate rate, *i.e.*, 0.1-10°C/s), an increase of the PQ clad ductility has been observed for Zircaloy-4 pre-hydrided at 600-700 wt.ppm [60]. This cooling scenario effect has been attributed to the hydrogen and oxygen micro-chemical partitioning occurring upon the on-cooling $\beta_{Zr} \rightarrow \alpha_{Zr}$ phase transformation, and to its consequences on the local prior- β_{Zr} hardening and embrittlement at RT, related to the intermediate cooling rate and the temperature from which the final water quenching is applied [62]. Thus, in the following, one has to keep in mind that the direct quenching conditions applied here are likely conservative compared to two-steps cooling. In other words, it is possible that the direct quenching applied induces higher PQ clad embrittlement compared to more LOCA-representative two-steps cooling.

- As already extensively used by M. Grosse *et al.* [41]-[46] [63] to study HT secondary hydriding, non-destructive neutron radiography/tomography has been performed to have a better view of the hydrogen redistribution, in three dimensions, in the samples that have experienced internal pressure-induced burst followed by HT oxidation and quenching. The experiments were performed on the IMAGINE radiography station connected to the experimental Orphée reactor at CEA Saclay. Compared to previous measurements [47], smaller spatial resolution was used (voxel size down to 50 μ m) to be able to map the hydrogen content not only as an integrated value for a given azimuthal position but also across the thickness of the cladding wall. The hydrogen quantity in the cladding was normalized with respect to reference samples with calibrated hydrogen contents. The typical sensitivity to hydrogen content is of the order of 100 wt.ppm.

- Additional PQ destructive hydrogen content measurements were performed by an Inert Gas Fusion Thermal Conductivity (IGFTC) technique using a *HORIBA EMGA-821 analyser*. Electron Probe Micro-Analysis (EPMA) using a *Cameca SX-100* microprobe was carried out to quantify the oxygen (and other alloying elements) concentration profile through the clad wall thickness.

- Local hardening of the quenched prior- β_{Zr} material is sensitive to the additional oxygen resulting from oxygen diffusion induced by the HT steam oxidation process. Thus, one can roughly evaluate the local oxygen content from micro-hardness measurements by using the following equation from [60].

$$HV(0.1) \text{ Prior-}\beta_{Zr} \text{ Zircaloy-4} = 370 \times [\text{wt.}\% \text{ O}] + 224 \quad \text{Equation 1}$$

where HV(0.1) denotes the Vickers hardness at 0.1 kg. Alternatively, the following relationship can be derived from the data already presented and discussed in [47]. This relationship is more representative of the hardening of the quenched prior- β_{Zr} microstructure of highly hydrided materials, *i.e.*, up to ~3000-4000 wt.ppm of hydrogen.

$$HV(0.1) \text{ Prior-}\beta_{Zr} \text{ Highly Hydrided-Zircaloy-4} = 193 \times [\text{wt.}\% \text{ O}] + 301 \quad \text{Equation 2}$$

Then, systematic PQ micro-hardness measurements have been carried out using a Buehler VH-3300 device equipped with Vickers or Knoop indenters. Up to 50 indents were performed throughout the clad wall thickness on each sample characterized after quenching, under an applied load of 0.05, 0.01 or 0.025 kg. These low load levels have been chosen to obtain a good description of the gradient through

the clad thickness. It has been checked that these low load levels gave results comparable to those obtained from Vickers hardness measurements performed at 0.1 kg, as more conventionally done.

- Finally, PQ RCTs have been carried out on 10 mm-long clad segments machined from the samples oxidized at HT. The tests were conducted on a universal electromechanical tensile machine at temperatures of 20 and 135°C (*saturation temperature during reflood in the case of large break LOCA*), with a cross-head displacement rate of 0.5 mm/min (*up to a maximum displacement of 6 mm or a maximum load of 2 kN*). Failure of RCT samples was considered to occur at the first significant load drop of more than 30%, which may indicate the formation of at least one through-wall crack. The fracture surface of the samples was examined using Scanning Electron Microscopy (SEM) to assess the PQ failure mode.

Figure 2-(b) shows the locations where the samples were machined from the tested clad segments for PQ analysis.

3. RESULTS AND DISCUSSION

3.1 As-received conditions

Zr-based substrate: As already mentioned, the “old-grade” HSn-HO-Zirc-4 material used here for the substrate is not fully representative of modern advanced nuclear fuel cladding made of zirconium-based alloys. Figure 3-(a) illustrates its as-received microstructure with a homogenous distribution of equiaxed α_{Zr} grains. The corresponding Vickers micro-hardness value is 200 HV_{0.05}, which is typical of fully recrystallized α_{Zr} microstructure. Additionally, a typical (non-oxidized) quenched prior- β_{Zr} microstructure is depicted in Figure 3-(b) showing a Vickers micro-hardness of 275 HV_{0.05}. Such microstructure has been obtained after a “flash” β_{Zr} thermal treatment at 1000°C (<30s) followed by water quenching down to RT to avoid significant HT oxidation. Thus, the hardness value measured seems to be higher than the typical one obtained for low-tin Zircaloy-4 prior- β_{Zr} (~220 HV_{0.05}). This is likely due to the higher oxygen and tin contents of HSn-HO-Zirc-4.

Outer coating: Figure 4 illustrates the typical as-received microstructure of the ~10 μ m-thick Cr outer-PVD-coated materials. Multiscale characterizations (down to the atomic scale) of the microstructures of such PVD-deposited Cr coating and Cr-Zr interface have been already reported and discussed in [11][20][64]. Just recall that, by using convenient PVD deposition parameters and process, such coatings are characterized by:

- low surface roughness and good thickness uniformity,
- very few defects/micro-cracks,
- no delamination or porosities at the Cr-Zr interface, thus inducing high bonding strength, even after (ion) irradiation [65],
- columnar grains morphology with specific crystallographic textures,
- reduced compressive internal stresses.

Inner coating: Detailed information about the typical microstructure of DLI-MOCVD Cr_xC_y inner coating has been already published in [26] and [27]. However, it is necessary to briefly discuss the particular microstructure of the inner-coated clad segment studied here. Figure 5 shows the typical aspect of the clad wall thickness at the “M2” axial location (Figure 1), corresponding to ~20 μ m-thick inner Cr_xC_y coating. One can observe the typical fully recrystallized structure of the HSn-HO-Zirc-4 substrate and a quite uniform inner coating thickness. Recall that previous studies have shown that such coating is in an amorphous state in as-received conditions.

However, localized delamination and porosities are observed at the Zr/Cr_xC_y interface. This could be partially related to the poor initial surface quality of the “old-grade” material used here as a substrate, as shown in Figure 6–(a). The SEM micrograph in Figure 6–(b) illustrates the typical morphology of the inner coating surface. The coating surface appears to have a quite smooth aspect with localized circular protrusions. Some localized cracks and delamination (*not illustrated here*) are also observed, indicating that an additional optimization of the deposition process and a substrate material with a smoother initial surface are necessary to further improve the inner coating adhesion properties.

Additionally, Figure 7 shows mapping and quantitative profiles of chemical elements obtained by EPMA within the wall thickness of the as-received 20 μ m-thick amorphous Cr_xC_y inner coating at the “M2” axial location. The average chemical stoichiometry of the coating deduced by integrating the EPMA profiles is close to Cr_{0.58}C_{0.30}O_{0.12}, which is in very good agreement with previous X-ray Photoelectron Spectroscopy (XPS) measurements reported in [26]. However, a significant oxygen content is observed, especially near the Cr_xC_y–Zr interface. Such an oxygen pollution may be attributed to the CVD process itself but also to the poor quality of the inner surface of the substrate (*i.e.*, with thin zirconium oxide and/or hydroxide scales initially present on its surface, before the CVD deposition process). Interestingly and neglecting the oxygen contribution, the chemical profiles in Figure 7–(b) show a continuous evolution of the Cr/C ratio with an apparent stoichiometry going from Cr₇C₃ near the Cr_xC_y–Zr interface to Cr₃C₂ close to the outer coating surface. Between these two zones, and based on previous studies and interpretation, one can make the assumption that there are nanometric C-free domains embedded in an amorphous Cr₇C₃ carbide matrix. Thus, in the following, this specific as-received amorphous nanocomposite structure and the existence of a gradient in Cr/C concentration ratio along the coating wall thickness have to be kept in mind, for further interpretation of the coating microstructural evolution upon HT steam oxidation, typical of LOCA situations.

3.2 HT two-sided steam oxidation and PQ RCT of inner/outer-coated short clad segments

Weight gain evolution and hydrogen uptake – The HT steam oxidation conditions tested and the associated weight gains and PQ hydrogen contents measured for uncoated and coated materials are summarized in Table 2. All the tested 30mm-long clad segments kept their integrity after the final direct water quenching down to RT, except the uncoated one oxidized for 3220s at 1200°C, consistently with its very high “Baker-Just” [66] “Equivalent Cladding Reacted” (ECR_{BJ}) calculated value of 70%.

The evolution of weight gain as a function of the two-sided oxidation time in steam at 1200°C is plotted in Figure 8 for uncoated and coated 30mm-long clad segments. The results are compared with one-sided steam oxidation data previously obtained at CEA on uncoated low-tin Zircaloy-4 [60] and on 12-15 μ m-thick Cr outer-coated M5_{Framatome} claddings [21]. From this figure, the following observations can be made:

- Despite its non-optimized chemical composition and poor inner surface quality, the old-grade HSn-HO-Zirc-4 behaves more or less like modern low-tin Zircaloy-4.
- The inner (10-20 μ m thick) DLI-MOCVD Cr_xC_y and/or the outer (~10 μ m-thick) PVD Cr seem to bring good protectiveness against steam oxidation at least up to 1500s (two-sided ECR_{BJ} = 47.5%) at 1200°C. Indeed, the weight gains of inner or outer one-sided coated materials are nearly half those of the uncoated materials after two-sided oxidation for 600-1500s. This means that, for these particular cases, the measured weight gain is essentially due to the oxidation of the (outer or inner) uncoated (and thus unprotected) clad surface. This is further confirmed by the behavior of the two-sided (inner + outer) coated samples, showing a very limited overall weight gain of only 2.2 mg/cm² after two-sided steam oxidation up to 1500s at 1200°C. As shown in Figure 8, this low weight gain value is comparable to the one achieved after one-sided (outer) steam oxidation for 1500s at 1200°C of 12-15 μ m-thick (PVD) Cr outer-coated M5_{Framatome} [21].
- For the higher (two-sided) oxidation time tested (3220s), the measured weight gain of the inner-coated sample is close to the uncoated reference one. This indicates that, for such a high ECR_{BJ} value (70%), the 10-20 μ m thick DLI-MOCVD Cr_xC_y coating is no more protective. Indeed, as already illustrated in [27], for such a long oxidation time, both uncoated and inner-coated samples have been almost fully transformed into zirconia and α_{Zr} (O) layers, thus inducing a high brittleness. However, it should be noticed that, for the same (one-sided) oxidation time at 1200°C, a 12-15 μ m-thick PVD Cr outer coating on a M5_{Framatome} substrate is still protective [21].
- Additionally, as shown in Table 2, the hydrogen contents measured post-oxidation show negligible or very limited hydrogen uptake, except for the inner-coated sample tested for 3220s at 1200°C for which the measured hydrogen concentration is close to 160 wt.ppm. Such phenomena could be due to a “transient” hydrogen uptake, as already observed on 12-15 μ m-thick (PVD) Cr outer-coated M5_{Framatome}

for equivalent ECR values (*i.e.* for one-sided ECR_{BJ} higher than 50% at 1200-1300°C) [20]. This has been attributed to the short period when the coating loses its protectiveness, thus inducing a direct access of steam molecules to the metallic zirconium-based substrate. For longer oxidation times, it was then observed that there was no more additional hydrogen uptake. It was thus assumed that, beyond this short period of transient hydrogen uptake, additional hydrogen uptake is prevented by the subsequent rapid growth of a dense and thick enough sub-coating zirconia layer. It would be interesting to check if such saturation of the hydrogen content is also operant for the present inner Cr_xC_y coating, when tested at longer oxidation times (*i.e.*, beyond the oxidation time when the coating loses its protectiveness).

In summary, based on the measured weight gains, one may conclude that, overall, the inner (10-20µm thick) DLI-MOCVD Cr_xC_y coating associated with an outer (~10µm thick) PVD Cr coating brings a significant coping time upon two-sided HT steam oxidation, up to an ECR_{BJ} value of ~50% (>1500s) at 1200°C. Such performances have been already achieved on PVD Cr-outer-coated materials, but in that case, only upon one-sided outer oxidation at HT [21].

Materials (substrate = HSn-HO-Zirc-4)	Initial outer / inner coating thickness (µm)	Two- sided steam oxidation time at 1200°C (s)	Calculated ECR_{BJ} (%) (*)	Measured weight gain (mg/cm ²)	Post- oxidation hydrogen content (wt.ppm)	Post- quenching integrity
Uncoated	0 / 0	600	30	14.88	27	YES
		3220	70	34.63	51	NO(**)
Cr_xC_y inner coated "O2"	0 / ~20	600	30	8.3	37	YES
Cr_xC_y inner coated "O1"	0 / ~20	3220	70	29.22	164	YES/NO (***)
Cr outer coated	9 / 0	600	30	7.56	25	YES
Cr outer coated	9 / 0	1500	47.5	11.8	31	YES
Cr/ Cr_xC_y two-sided-coated "O4"	9 / 9	600	30	1.05	35	YES
Cr/ Cr_xC_y two-sided-coated "O3"	9 / 11	1500	47.5	2.22	48	YES

Table 2 - HT steam oxidation conditions, associated weight gains and PQ hydrogen contents of uncoated, and of inner and/or outer-coated HSn-HO-Zirc-4 (30mm-long) clad segments

(*) Equivalent Cladding Reacted (ECR) parameter calculated using the "Baker-Just" correlation for two-sided oxidation of uncoated 610µm-thick cladding tubes

(**) longitudinal (through-wall) crack; high PQ embrittlement (failure upon machining/cutting)

(***) external rough protrusions, but with no apparent through-wall cracking; high PQ embrittlement (failure upon machining/cutting)

Cr_xC_y inner coating microstructure after steam oxidation for 600s at 1200°C – Post-HT-oxidation microstructures of both outer Cr PVD and inner DLI-MOCVD Cr_xC_y coatings have been already described in details in [11][20][21] and **Erreur ! Source du renvoi introuvable.**[27], respectively. However, it seems useful to recall some Electron-Back Scattered Diffraction (EBSD) results obtained on the inner Cr_xC_y coating, after two-sided steam oxidation for 600s at 1200°C, as illustrated in Figure 9 (replotted from [27]). In this figure, one can observe that preferential recrystallization and formation of Cr_7C_3 and Cr_3C_2 have occurred in the vicinities of the Cr_xC_y -Zr interface and the outer coating surface, respectively. This can be understood from the EPMA results already shown in Figure 7. Indeed, their localization coincides with the coating depth where the Cr/C atomic fraction ratio is close to the stoichiometries of carbides, thus promoting their crystallization upon HT incursion. By contrast, at the coating mid-thickness, EBSD analysis failed to identify recrystallized carbides phases, while submicronic chromia crystallites have been identified. These last ones are likely due to the initial oxygen content (*i.e.*, ~10 atomic%) within the amorphous Cr_xC_y coating layer and to potential additional oxygen diffusion from the outer oxidized coating surface which took place upon steam oxidation at 1200°C. However, at coating mid-thickness, one can assume that the observed delay in carbides crystallization

could be due to the local as-received particular nanocomposite structure, with some nanometric carbon-free domains embedded in an amorphous Cr_7C_3 carbide matrix [27]. It is possible that such a non-stoichiometric as-received (amorphous) microstructure, with some carbon-free submicronic areas, is less favorable to further crystallization of HT chromium carbides. As already discussed in [27], maintaining such an amorphous (crystallite free) microstructure as long as possible at HT could delay the overall oxidation kinetics of the coating, by decreasing the density of grain boundaries which can be assumed to be accelerated diffusion paths for oxygen. Additionally, in this figure, one can observe a more or less continuous, micrometer-thick, zirconium carbide layer at the Cr_xC_y -Zr interface, which has been induced by (spatially limited) Zr-C interdiffusion phenomena at HT.

Effects of pre-cracks/defects in the Cr_xC_y coating – One important issue of coated claddings, especially upon LOCA, is the consequence of potential defects/cracks in the coating on the subsequent (localized) HT oxidation and the effect on the cladding residual strength and ductility upon quenching and post-quenching situations. This is one of the fundamental reason for having chosen pure metallic chromium coating for the outer clad protection instead of ceramic ones, during the early studies carried out at CEA on several types of coatings [11]. Indeed, it has been shown that, upon LOCA transient, optimized PVD Cr metallic coating is able to sustain significant clad ballooning without cracking or spallation, thus preserving the protection against the subsequent HT oxidation, even at the balloon location [67]-[69]. Because of the amorphous and/or ceramic nature of the inner Cr_xC_y coating studied here, it is anticipated that this type of coating may experience significant cracking upon clad deformation (ballooning) typical of LOCA situations. Figure 10 depicts typical zones in samples with an inner Cr_xC_y coating with pre-existing cracks that have experienced HT steam oxidation and quenching. One can observe three typical (localized) oxidation behaviors at the locations of the defects/cracks in the inner coating:

- *Type-1 coating crack/defect with no oxygen diffusion into the Zr substrate.* This particular situation seems to be due to the coating self-healing capacity, *i.e.*, formation of a dense and thick enough Cr_2O_3 scale inside the cracking, thus preserving the Zr substrate from further oxygen penetration and diffusion.
- *Type-2 coating crack/defect with partial oxygen diffusion into the Zr-based substrate inducing $\alpha_{\text{Zr}}(\text{O})$ incursions (partial self-healing capacity).* Such a situation may occur when the inward oxygen diffusion flux into the zirconium substrate is not high enough to promote zirconium oxide formation but sufficient to induce $\alpha_{\text{Zr}}(\text{O})$ formation. In fact, such a situation has also been observed, transitorily, for the outer PVD Cr-coated cladding when the Cr coating begins to be no more protective, upon long enough HT oxidation [20][21].
- *Type-3 coating crack/defect with under-coating zirconia spot formation (no self-healing).* In that particular case, the rapid formation of under-coating zirconia, due to a direct access of steam to the Zr-based substrate, leads to opening of the pre-existing crack/defect due to the associated volume change (*i.e.*, “Pilling-Bedworth” ratio). This situation has been already observed on outer PVD Cr-coated cladding when the coating loses definitely its HT oxidation protectiveness [20][21]. However, as already observed on PVD outer Cr-coated Zr-based claddings, there is no sign of local oxidation acceleration, nor of extended coating delamination and spallation at the vicinity of the cracks. Indeed, the freshly formed sub-coating zirconia spot is always thinner (or at least equivalent) than the zirconia scale formed in the same HT oxidation conditions on uncoated Zr-cladding, as already observed for PVD coated materials [11][70][75].

It must be pointed out that such defects were not so numerous on the (not deformed) clad segments tested. It is why they did not significantly affect the overall oxidation kinetics as derived from the measured weight gain evolution. However, after significant clad deformation (ballooning), the surface density of such through-wall cracks in the coating may increase and thus may affect the PQ strength and ductility of inner Cr_xC_y coated materials that have experienced ballooning and burst, followed by HT steam oxidation of the inner (coated) clad surface.

EPMA and micro-hardness profiles - As initially reported by Sawatzky [71] and Chung & Kassner [72], oxygen content in the residual prior- β_{Zr} layer is a key parameter to predict the post-quench mechanical behavior of as-received zirconium-based nuclear fuel cladding tubes, after HT steam oxidation and final water quenching typical of LOCA situations. More recently, the critical oxygen content for water-quenched prior- β_{Zr} structure has been reassessed for both Zircaloy-4 and Zr-Nb alloys [60][60][73][74].

For example, based on PQ impact tests, a critical oxygen content value of 0.4 +/- 0.1 wt.% has been related to the Ductile-to-Brittle (DBT) failure mode transition of (as-water-quenched) prior- β_{Zr} at RT, for both Zircaloy-4 and M5_{Framatome} alloys [60]. Then, in the following, particular attention is paid to the PQ oxygen (but also chromium) diffusion profiles through the cladding thickness measured by EPMA, especially in the residual prior- β_{Zr} layer. Indeed, this last layer is the only residual part of the cladding able to keep significant ductility and toughness after HT steam oxidation and quenching. Then, the oxygen (and chromium) diffusion profiles can be related to the associated hardening of the as-quenched prior- β_{Zr} structure. For this purpose, Table 3 and Figure 11 to Figure 16 summarize the oxygen and chromium diffusion profiles measured by EPMA and the associated micro-hardness profiles for the different types of coated samples, after two-sided steam oxidation up to 1500s at 1200°C (*i.e.*, up to an ECR_{BJ} value of nearly 50%). The two uncoated / inner-coated clad segments oxidized up to 3220s have not been considered here because, due to the very high ECR_{BJ} achieved (70%), these samples have been fully transformed into brittle zirconia and $\alpha_{Zr}(O)$ thick layers, as already shown and discussed in [27]. One can make the following observations:

- *After two-sided oxidation for 600s at 1200°C (ECR_{BJ} = 30%):* due to the quite high ECR achieved, the uncoated clad is characterized by a very limited residual prior- β_{Zr} layer thickness, with an oxygen content of ~0.6 wt.%. Such an oxygen concentration is consistent with the (maximum) oxygen solubility in the β_{Zr} phase at 1200°C which can be derived from the pseudo-binary oxygen-Zircaloy-4 phase diagram proposed by Chung & Kassner (*i.e.*, 0.57 wt.%) [76]. This oxygen concentration is also consistent with the measurements already performed at CEA on low-tin Zircaloy-4 one-sided steam oxidized at 1200°C [60] and calculated using Thermocalc® software associated with the CEA “Zircobase” thermodynamic database for zirconium alloys [77]. In comparison, both one-sided and two-sided coated materials are characterized by thicker residual prior- β_{Zr} layers with lower oxygen contents due to protection from oxygen ingress provided by the inner and/or outer coating. However, probably due to surface oxidation contamination effects, it seems that EPMA measurements have overestimated the prior- β_{Zr} oxygen content of the two-sided-coated materials when compared to the value which can be derived from the micro-hardness measurements (using equation 1). Then, for this particular sample and by taking into account the oxygen content derived from the measured prior- β_{Zr} hardening, one can conclude that the two-sided coatings have reduced drastically the oxygen diffusion within the metallic zirconium-based substrate upon the HT two-sided steam oxidation.

- *After two-sided oxidation for 1500s at 1200°C (ECR_{BJ} = 47.5%):* for such a high ECR, the one-sided outer-coated materials have experienced significant oxygen diffusion from the unprotected inner surface, inducing deep $\alpha_{Zr}(O)$ incursions. Overall, one can consider that until the Cr coating is protective, two-sided oxidation of one-sided (outer) coated materials is more or less comparable to one-sided oxidation of uncoated materials, as also reflected by their overall weight gain. For two-sided coated materials, one can observe that the inner coating has been more or less protective while slight oxygen diffusion has been observed at the outer Cr-coated side. Based on the HT steam oxidation mechanism and process of outer (PVD) Cr-coated Zr-based claddings already discussed in [20], this particular oxidation time at 1200°C may coincide with the early beginning of oxygen intergranular diffusion through the residual metallic (sub-outer-chromia scale) Cr coating thickness. Furthermore, the Cr-coating consumption kinetics already described in [11] predict that 9 μ m-thick Cr coating may be fully consumed after nearly 3000s at 1200°C, which seems to be consistent with the results obtained here.

Finally, as already reported in [11], it can be observed that, upon HT incursion, Cr diffusion occurred within the β_{Zr} substrate from the Cr-coated outer surface, while no apparent Cr diffusion is observed from the Cr_xC_y inner-coated surface. Indeed and as already mentioned in [27], for short oxidation times, only slight carbon diffusion has been detected by EPMA from the Cr_xC_y inner-coated surface (*not represented in the figures for sake of simplicity*). Additionally, for the coated samples oxidized for 600s, it can be observed that, on the outer Cr-coated side, chromium diffusion has induced a significant PQ prior- β_{Zr} hardening, while no hardening is observed at the vicinity of the Cr_xC_y inner-coated surface.

Materials (HSn-HO-Zirc-4 substrate)	two-sided oxidation time (s) at 1200°C / ECR_{BJ} (%)	wt.% O in prior- β_{Zr} (EPMA)	HV(0.01)	wt.% O in prior- β_{Zr} (calculated from HV, eq. 1)
Uncoated	600 / 30	0.60 +/- 0.10	<i>not measured</i>	0.60 (from EPMA)
Cr _x C _y one-sided inner-coated	600 / 30	0.44 +/- 0.10	370 +/- 10	0.39
Cr one-sided outer-coated	600 / 30	0.48 +/- 0.10	405 +/- 10	0.49
Cr one-sided outer-coated	1500 / 47.5	~1.1	~475	~0.7
Cr/Cr _x C _y two-sided coated "O4"	600 / 30	< 0.4 (*)	300 +/- 10	0.21
Cr/Cr _x C _y two-sided coated "O3"	1500 / 47.5	< 0.6 (*)	350 +/- 10	0.34

(*) likely overestimated values by EPMA, due to surface contamination/oxidation

Table 3 - Evaluation of the oxygen contents of residual prior- β_{Zr} layer from EPMA measurements and calculated from micro-hardness using equation 1

PQ RCT – Preliminary remark: The determination of the “Residual PQ clad ductility” parameter from the RCT results was sometimes not obvious. Indeed, especially for the two-sided coated materials, it appeared that several RCT curves show multiple significant load drops, but with a relative amplitude lower than 30%, which is the threshold value that is conventionally considered to be representative of through-wall clad cracking upon PQ RCT of reference (uncoated) zirconium-based claddings ([30][80][81]). Such PQ RCT specific behaviors may be related to the more complex multilayer PQ microstructure of HT oxidized inner-outer coated materials, compared to reference uncoated claddings. Complementary experiments are on-going at CEA to better assess the complex mechanical behavior of multilayered materials following HT oxidation and quenching, and will need further analysis and discussion. Keeping that in mind, in the following, it was decided to consider only the maximum load measured during the RCT tests, which is assumed to be a less ambiguous parameter than the “residual ductility” one, for the particular samples tested here.

“Qualitative” evaluations of the overall PQ failure mode and maximum load values derived from PQ RCTs are presented in Table 4 and plotted as a function of the ECR_{BJ} values (calculated for two-sided oxidation conditions) in Figure 17. They are compared to previous data obtained on reference uncoated [60] [79] and outer Cr-coated [21] cladding materials. Overall, it is interesting to note that:

- when plotted as a function of calculated ECR_{BJ} , the residual strength of the two-sided coated claddings that have experienced HT two-sided steam oxidation and direct quenching is close to the one obtained for outer-Cr-coated M5_{Framatome} materials, but in this last case after one-sided (outer) oxidation corresponding to oxidation times four times longer;
- one-sided (inner and outer) coated materials behave quite similarly and display an intermediate PQ “residual strength” evolution as a function of the calculated ECR_{BJ} , ranging between the respective ones of uncoated and two-sided coated materials.

Materials (HSn-HO-Zirc-4 substrate)	two-sided oxidation time (s) at 1200°C / ECR _{BJ} (%)	Room Temperature		135°C	
		Overall failure mode (*)	Max. load (N)	Overall failure mode (*)	Max. load (N)
Uncoated	600 / 30	Brittle	225	Low ductility	486
	1500 / 47.5	Failed upon Q or PQ machining			
Cr _x C _y one-sided inner-coated "O2"	600 / 30	Low ductility	486	Low ductility	498
Cr _x C _y one-sided inner-coated "O1"	1500 / 47.5	Failed upon Q or PQ machining			
Cr one-sided outer-coated	600 / 30	Low ductility	387	Low ductility	482
Cr one-sided outer-coated	1500 / 47.5	Brittle	165	Brittle	226
Cr/Cr _x C _y 2-sides coated "O4"	600 / 30	Fully Ductile	1048	Fully Ductile	1098
Cr/Cr _x C _y 2-sides coated "O3"	1500 / 47.5	Low ductility	545	Low ductility	371

(*) Qualitative assessment, in some cases, "low ductility" should mean "macroscopically nearly brittle"

Table 4 – Maximum load derived from PQ RCTs obtained after two-sided steam oxidation at 1200°C followed by direct quenching on inner and/or outer-coated 1cm-long clad segments

Typical fractographs showing the failure mode of the prior- β_{Zr} layer at RT, corresponding to clad mid-thickness location, are presented in Figure 18. These observations are consistent with the qualitative overall failure mode given in Table 4. Thus, it can be observed that the DBT at RT of uncoated claddings occurs at an ECR_{BJ} value close to (or lower than) 30%, while one-sided coated materials show a DBT ranging between ECR_{BJ} values of 30 and 47% and two-sided coated materials seem to keep significant residual ductility up to at least an ECR_{BJ} value of 47%.

So, as far as the calculated ECR parameter is considered, it can be concluded that the two-sided coated cladding prototype materials studied here show comparable PQ residual mechanical strength after two-sided steam oxidation (at 1200°C) when compared to reference (outer) Cr-coated M5_{Framatome} materials after one-sided (outer) HT oxidation. In other words, the 10 μ m-thick Cr_xC_y inner coating studied here brings comparable protectiveness against HT steam oxidation when compared to outer PVD chromium coatings with comparable thicknesses.

Finally and as previously mentioned, PQ mechanical properties of zirconium-based nuclear fuel claddings that have experienced HT steam oxidation and a final water quenching mainly rely on the residual prior- β_{Zr} layer thickness and its oxygen content [60][71]-[74]. To address this effect, Figure 19 illustrates the evolution of the maximum load derived from PQ RCTs as a function of the prior- β_{Zr} oxygen content (*derived from micro-hardness measurements - equation 1*). The observed PQ "residual strength" and the associated evolution of the residual ductility of HT oxidized inner and/or outer coated Zircaloy-4 appear to be consistent with previous data obtained on uncoated [60][79] and outer Cr-coated [12][21] cladding materials.

3.3 Inner steam oxidation and secondary hydriding following clad ballooning and burst, and consequences on the residual strength derived from RCT

Let us recall that four tests consisting in a first temperature ramp test under internal pressure up to burst, followed by steam oxidation at 1200°C and final direct water quenching, have been carried out on both uncoated and inner-coated HSn-HO-Zirc-4 clad segments (*with internal alumina rods to simulate the gap between clad inner surface and fuel oxide pellets*). The tests conditions and the associated measured burst temperatures and circumferential strains are given in Table 5. As already observed for outer Cr-coated low-tin Zircaloy-4 and M5_{Framatome} claddings [67]-[69], the presence of the coating tends

to decrease clad ballooning when compared to uncoated materials tested in the same conditions. Let us recall that such behavior may have positive consequences on flow blockage and potential fuel pellets fragments relocalisation in the balloon upon LOCA transient. Moreover, the thicker inner coating tested (*i.e.*, with a thickness ranging between 11 and 19 μ m) has induced a significant increase of the burst temperature, indicating that such inner coating may have a clad strengthening effect at HT, as already observed for outer 12-15 μ m Cr-coated materials [67]-[69].

Materials	Internal pressure "EDGAR" thermal ramp tests					HT steam oxidation tests	
	Heating rate (°C/s)	Internal pressure (bar)	Burst temperature (°C)	Total hoop strain at burst location (%)	Hoop strain at +/-20mm from burst location (%)	Steam oxidation time at 1200°C (s)	ECR _{BJ} (%)
Uncoated HSn-HO-Zirc-4	25	40	981	13	3.5	600	30 - 35(*)
11-19 μ m thick Cr _x C _y inner coated HSn-HO-Zirc-4	25	40	1045	6,3	0.7	600	30 - 32(*)
Uncoated HSn-HO-Zirc-4	25	40	979	13,4	2.7	1500	47.5 - 55(*)
6-9 μ m thick Cr _x C _y inner coated HSn-HO-Zirc-4	25	40	977	4,6	1.8	1500	47.5 - 50(*)

(*) Max. ECR_{BJ} values calculated by taking into account the wall clad thinning at burst location

Table 5 - Internal pressure temperature ramp tests and associated burst temperatures and mean circumferential strains, and corresponding subsequent HT steam oxidation test conditions followed by direct water quenching

Figure 20 depicts the macroscopic aspects of the four uncoated and inner-coated HSn-HO-Zirc-4 clad segments that have experienced ballooning and burst tests then subsequent HT steam oxidation and direct water quenching. After oxidation for 600s at 1200°C corresponding to an ECR_{BJ} value of at least 30%, the inner-coated clad survived to the final quenching while the uncoated materials did not. After oxidation for 1500s at 1200°C (ECR_{BJ} of at least 47.5%), both materials failed upon quenching but the uncoated one has broken in much more fragments than the inner-coated one. For this last one it must be also kept in mind that the initial inner coating thickness was limited to 6-9 μ m, thus likely inducing a lower protectiveness of the inner clad surface upon the HT steam oxidation step, when compared to the one tested for 600s at 1200°C, with inner coating thickness ranging from 11 to 19 μ m. However, these preliminary macroscopic results show encouraging behavior of inner-coated claddings, with indications of a significant additional coping time before failure upon quenching, when compared to reference uncoated materials.

In the following, we will focus on samples tested for 600s at 1200°C (ECR_{BJ} = 30 – 35%) only, because multi-fragmentation of clad segments that have experienced HT steam oxidation up to ECR_{BJ} values of nearly 50% makes more complicated the accurate PQ metallurgical examinations and hydrogen measurements as a function of the PQ clad segment location. Additionally, for such fragmented PQ clad segments, it was obviously not possible to perform PQ RCTs.

PQ microstructures, phase thicknesses - Figure 21 shows typical optical micrographs of the PQ uncoated and inner-coated microstructures of HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst temperature ramp tests then subsequent HT steam oxidation for 600s at 1200°C followed by direct water quenching down to RT, for different axial locations from the burst. The corresponding oxide and metallic sub-layers thicknesses are plotted in Figure 22. In this last figure, the results are compared to the results of previous tests carried out on (uncoated) M5_{Framatome} tested using the same methodology, but for a shorter HT oxidation time at 1200°C, that is 160s, which corresponds to a two-sided ECR_{BJ} value of nearly 17% [47].

One can make the following observations:

- *For the outer part of the claddings* : The respective thicknesses of the ZrO_2 and $\alpha_{Zr}(O)$ layers are very close to the ones already measured on low-tin Zircaloy-4 after one-sided (outer) steam oxidation for 600s at 1200°C [60], thus confirming the consistency and the good reproducibility of the HT steam oxidation tests performed so far on uncoated reference materials. Moreover, the ZrO_2 and $\alpha_{Zr}(O)$ thicknesses achieved after 600s at 1200°C are twice the values measured on (uncoated) tested M5_{Framatome} after 160s at 1200°C [47]. This is fully consistent with the ECR_{BJ} value here achieved, after 600s of oxidation, being twice the value achieved after 160s oxidation, thanks to the parabolic oxidation kinetics of the materials when tested at 1200°C.

- *For the inner part of the claddings* : On the one hand, the thicknesses of the ZrO_2 and $\alpha_{Zr}(O)$ layers in uncoated HSn-HO-Zirc-4 oxidized for 600s at 1200°C are comparable to those of M5_{Framatome} oxidized for 160s at 1200°C, with, close to the burst, comparable values to the ones measured at the outer surface. This is likely due to the easy steam access through the burst opening inducing locally two-sided oxidation conditions. Close to the sample ends and as already discussed in details in [47], there is no more inner oxide scale and the $\alpha_{Zr}(O)$ sub-layer is no more circumferentially homogeneous (Figure 21). This can be related to steam starvation conditions that prevail within the gap between the inner rod (simulating fuels pellets) and the inner clad surface, close to the end caps. Such a behavior is believed to be responsible for the localized massive (secondary) hydrogen uptake, due to: (i) gaseous hydrogen enrichment (and associated oxygen depletion) in the gap and (ii) the absence of continuous and dense zirconia scale which could acts as an efficient hydrogen permeation barrier.

For the inner-coated clad segment, most of the inner surface has been protected from significant oxidation and oxygen ingress. However, localized $\alpha_{Zr}(O)$ areas can be observed (Figure 21-b), likely due to localized inner coating cracking and/or spallation. It is worth noting that the localized $\alpha_{Zr}(O)$ growth is preferentially observed at the vicinity of the burst opening. This is likely due to higher localized strain promoting steam/oxygen access due to larger (localized) inner coating cracks opening and/or density.

Secondary hydriding and associated quenched prior- β_{Zr} hardening – Figure 23 depicts the hydrogen distribution derived from non-destructive neutron tomography and from *post-mortem* IGFTC measurements. It is worth noticing that the two types of hydrogen measurements give fully consistent results. Consistently with the numerous previous studies dealing with secondary hydriding phenomena upon LOCA [30]-[52], the uncoated clad segment shows typical secondary hydrogen peaks with maximum values reaching ~2500 wt.ppm, close to the sample ends. While, for the inner-coated clad segment, the uptake of hydrogen appears to be much lower, thus demonstrating the overall capacity of the inner DLI-MOCVD Cr_xC_y coating to prevent from rapid hydrogen uptake (from the inner clad surface).

However, Figure 23-(b) shows an apparent axial gradient of the hydrogen content within the PQ inner-coated clad segment. In other words, upon HT inner clad oxidation, it appears that the inner coating protectiveness has not be the same at each end of the tested clad segment. Such behavior should be due to:

(i) the axial gradient in thickness of the as-received Cr_xC_y inner coating, decreasing from ~19 μ m to ~11 μ m, with a lower HT oxidation/hydriding protectiveness of the thinner part of the inner coating at one end of the clad segment;

(ii) larger local “damage” of the inner coating (*i.e.*, *higher surface density of cracks and/or spallation inducing locally an easier steam/hydrogen access to the Zircaloy substrate*) at one end of the PQ clad segment, as qualitatively observed in Figure 21-(b) in some parts of the PQ clad segment.

However, at that point, it is not possible to conclude if the Cr_xC_y coating acts as an intrinsic efficient hydrogen permeation barrier and/or if the lower hydrogen uptake observed is simply due to the lower generation of gaseous hydrogen within the gap, related to the overall slower oxidation of the inner clad surface.

Additionally, through-thickness micro-hardness profiles measured at different axial locations of the two clad segments tested are plotted in Figure 24. One can observe in the mid part of the claddings corresponding to the residual prior- β_{Zr} layer that the PQ hardening is significantly higher within the uncoated materials than in the inner-coated one. As already discussed in [47] and recalled below, this is likely due to an indirect effect of the hydrogen uptake, leading to an increase of the PQ prior- β_{Zr} oxygen content. Indeed, local PQ hydrogen contents, prior- β_{Zr} micro-hardness and associated estimated oxygen contents are listed in Table 6 as a function of the axial distance from the burst. Then, the oxygen content

derived from prior- β_{Zr} micro-hardness can be compared to “Thermocal + Zircobase” [84][85] predicted values (*assuming local thermodynamic equilibrium*). It is worth noticing that, even if the thermodynamic calculations sometimes overestimate the oxygen value (*potentially due to not fully at equilibrium conditions*), the present results and calculations confirm the thermodynamic hydrogen effect, inducing a significant increase of the oxygen solubility in the HT β_{Zr} phase, and thus, of the PQ prior- β_{Zr} hardness, as already discussed in more details in [47] and [60].

Axial distance from burst location (mm)	Wt.ppm H (from IGFTC measurements)		Mid-wall clad prior- β HV(0.01) (+ standard deviation)		Prior- β oxygen content derived from HV(0.01) using eq. 2 (wt.%)		Max. prior- β oxygen content from thermodynamic calculations at 1200°C vs. measured hydrogen content (averaged value in wt.%)	
	Uncoated	Inner coated	Uncoated	Inner coated	Uncoated	Inner coated	Uncoated	Inner coated
-65	2099	219	640 (45)	405 (45)	1.76	0.54	2.03	0.61
	2148	192						
	2303	201						
-40	2539	230	525 (45)	420 (45)	1.16	0.62	1.78	0.63
	1726	218						
	1420	231						
-15	51	209	615 (40) (*)	405 (45)	1.63 (*)	0.54	0.53	0.61
	58	178						
	49	223						
15	42	347	470 (50)	480 (27)	0.88	0.93	0.53	0.69
	36	410						
	38	290						
40	2483	387	490 (45)	445 (35)	0.98	0.75	2.01	0.76
	1899	491						
	2089	520						
65	2738	1008	570 (50)	470 (25)	1.39	0.88	2.25	1.02
	not measured	775						
	not measured	847						

(*) not fully representative prior- β_{Zr} values due to localized clad thinning inducing additional hardening due to $\alpha_{Zr}(O)$ incursions

Table 6 - PQ hydrogen contents, prior- β_{Zr} micro-hardness and associated estimated oxygen contents compared to “Thermocalc + Zircobase” predicted values, of uncoated and inner coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst temperature ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C + direct water quenching down to RT

PQ RCT behaviors – Table 7 gives the maximum loads derived from PQ RCTs carried out at 135°C on uncoated and inner-coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst tests, then HT steam oxidation for 600s at 1200°C. These values correspond to different axial locations of the tested rings along the PQ clad segments. Even if, for all the ring samples tested, the overall RCT behavior seems to be macroscopically nearly brittle with very limited or negligible residual ductility, the inner-coated materials show higher strength values. For the uncoated one, the samples located at the vicinity of the clad segment ends display negligible residual strength. In this last case, this can be related to the localized high hydrogen and associated oxygen contents, typical of secondary hydrogen peaks.

		Max load from RCTs at 135°C (N)			
		approximate distance from burst opening (mm)			
	ECR _{BJ} (%) (*)	-57 to -47	-22 to -32	+22 to +32	+47 to +57
Uncoated	30-35	~20 (**)	183	183	<20 (***)
Inner-coated	30-32	287	296	258	155

(*) ECR_{BJ} range estimated by taking into account the localized clad thinning due to ballooning

(**) failed upon RCT pre-loading

(***) failed upon machining

Table 7 - Maximum load derived from PQ RCTs carried out on 1cm-long ring samples at 135°C of uncoated and inner coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst thermal ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C

Figure 25 is an attempt to relate the residual PQ clad segments strength derived from RCT to the localized (circumferentially averaged) hydrogen and oxygen contents. As already discussed in [47] and as can be observed in Table 6, the localized prior- β_{Zr} oxygen and hydrogen concentrations are more or less linearly correlated. It is thus not surprising that plotting the strength measured during RCT vs. the concentration of each type of chemical elements gives the same overall trend. One can observe that beyond respectively 2000-3000 wt.ppm of hydrogen corresponding to oxygen contents ranging between ~1.1 and ~1.4 wt.%, the quenched prior- β_{Zr} structure is fully brittle, with negligible residual strength. Such hydrogen/oxygen concentrations in prior- β_{Zr} are believed to correspond to a fully brittle PQ behavior. Moreover, in Figure 25-(a), the PQ strength measured here from RCT are compared with some previous IRSN PQ tensile data [82] obtained on pre-hydrided Zircaloy-4 samples that have been two-sided steam oxidized at 1200°C up to an ECR_{BJ}=16%, and water quenched. Even if the PQ mechanical tests are not the same, the results seem to be quite consistent.

Finally, to go further, let us consider the recent work carried out by Le Hong *et al.* by using Zircaloy-4 clad samples homogeneously pre-charged in hydrogen and/or oxygen [86] [87]. Among other results, after applying HT incursion in the β_{Zr} temperature range and then cooling down to RT, the authors were able to derive laws for the produced prior- β_{Zr} “model structures”, describing the evolution of mechanical data as a function of both oxygen and hydrogen contents in β_{Zr} . For PQ macroscopically “brittle” materials, Le Hong *et al.* derived the equation below (eq. 3), which can be used to predict the residual PQ Ultimate Tensile Strength (UTS) in uniaxial tension of the prior- β_{Zr} structure at low temperature (here at RT), as a function of both the oxygen and the hydrogen concentrations, up to thousands of wt.ppm:

$$UTS_{Brittle} (MPa) = 684.0 \times (1 - 1.329 \cdot 10^{-4} C_H) \times (1 - 0.760 C_O) \quad \text{equation 3}$$

where C_H and C_O correspond to prior- β_{Zr} hydrogen concentration in wt.ppm and oxygen concentration in wt.%, respectively.

The PQ tensile strength values calculated *using eq. 3* are plotted in Figure 26 as a function of the experimental ones derived from PQ RCTs. Both two-sided oxidized short clad segments (with quite negligible hydrogen pick-up) and the two longer clad segments that have experienced steam oxidation following burst occurrence (with localized high hydrogen contents) have been considered. It can be concluded that, even if the mechanical tests are not the same, Le Hong *et al.*’ correlation gives a reasonable prediction of the PQ residual strength and confirm that both oxygen and hydrogen contribute to the post-LOCA localized embrittlement of the fuel cladding.

4. CONCLUSIONS AND FURTHER WORK

To further improve the properties of coated claddings as E-ATF, DLI-MOCVD Cr_xC_y coating has been developed and successfully deposited onto the inner surface of an old-grade high-tin Zircaloy-4 cladding tube prototype. Then, one-sided and two-sided steam oxidation tests have been carried out at 1200°C on outer PVD Cr-coated and/or inner Cr_xC_y -coated clad segments, with a final direct water quenching down to RT. Some of the HT oxidation tests have been performed after a first internal pressure-induced burst step to address inner oxidation and associated secondary hydriding, typical of (semi-)integral LOCA transients.

Among other results, it is shown that the inner 10-20µm-thick Cr_xC_y coatings bring protectiveness against steam oxidation at 1200°C comparable to that of outer PVD 10-15µm-thick Cr coating, as well as a significant reduction of secondary hydriding after clad ballooning and burst. Then, when compared to the reference uncoated claddings, DLI-MOCVD Cr_xC_y inner coated materials show a higher strength, upon and after final water quenching, related to lower oxygen and/or hydrogen uptake in the zirconium substrate, at least up to an ECR_{BJ} value of 30%.

Further experiments on advanced nuclear zirconium-based fuel cladding two-sided coated are planned, to confirm these preliminary encouraging results. In addition, it is planned to assess the performances of the inner coating under more representative in-service conditions and for more demanding accidental situations, including for example pellet-cladding mechanical interaction issues and chemical compatibility with the uranium oxide fuel pellets at HT.

ACKNOWLEDGEMENTS

Special thanks to:

- R. Boichot and M. Pons from SIMAP-CNRS (Grenoble, France) for their inputs into the R&D and fabrication of DLI-MOCVD inner clad coatings;
- Y. Gazal from CIRIMAT (Toulouse, France) for the growth of CrC_x inner coatings using a DLI-MOCVD pilot;
- E. Monsifrot and J. Runser from DEPHIS society (Etupes, France) for PVD (outer) Cr-coated materials fabrication;
- D. Hamon and A. Lequien from CEA/SRMA/LA2M, for the EPMA measurements and C. Miton for PQ hydrogen (IGFTC) analysis;
- J. Henry, L. Vincent and L. Portier from CEA/SRMA and H. Palancher from CEA/DEC for their advising and organizing support.

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(b) $HV_{0.05} = 275 \pm 5$

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Figure 9 - SEM-EBSD map in phase contrast imaging mode of the inner Cr_xCr_y coating, after steam oxidation for 600s at 1200°C, replotted from [27]

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(b) Type-2 coating crack/defect with partial oxygen diffusion into the Zr substrate inducing $\alpha_{\text{Zr}}(\text{O})$ incursions into the Zr-based substrate (partial self-healing capacity)

(c) Type-3 coating crack/defect with under-coating zirconia spot formation (no self-healing)

(*) Outer Ni layer deposited before sample polishing to preserve the oxide and coating scales

Figure 10 - EPMA chemical element mappings (a), optical micrograph (b) and SEM-BSE electron micrograph (c) of different types of defects/cracks in the inner DLI-MOCVD Cr_xC_y coating, as locally observed after steam oxidation for 600s at 1200°C

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Figure 13 - Optical micrograph and through-thickness (from the outer diameter) oxygen (and chromium) diffusion profiles from EPMA and μ -hardness profiles of outer-coated HSn-HO-Zirc-4 after two-sided steam oxidation at 1200°C for 600s

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Figure 17 – Maximum load values derived from PQ RCTs after steam oxidation at 1200°C followed by direct quenching down to RT (on 1cm-long clad segments), plotted as a function of the Baker-Just ECR and compared with previous data obtained on uncoated [60] [79] and outer Cr-coated [21] cladding materials

Figure 18 – Typical fractographs showing the prior- β_{Zr} failure mode at mid clad thickness location, after PQ RCTs carried out at RT

(a)

(b)

Figure 19 – Maximum load values derived from PQ RCTs after steam oxidation at 1200°C followed by direct quenching down to RT, plotted as a function of prior- β_{Zr} oxygen content derived from micro-hardness measurements (equation 1) and compared with previous data obtained on uncoated Low-Tin Zirc-4 [60] [79] and outer Cr-coated M5_{Framatome} [21] cladding materials

Remark: due to the RCT maximum displacement available, the points corresponding to 2000N are underestimated values (i.e., corresponding to un-failed samples)

Figure 20 - Visual aspects of uncoated and inner-coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst temperature ramp tests and then subsequent HT steam oxidation and direct water quenching down to RT

(a)

(b)

Figure 21 - Optical micrograph showing the PQ microstructures of uncoated (a) and inner-coated (b) HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst thermal pressure ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C

Figure 22 - Circumferentially averaged values of the thicknesses of the inner/outer ZrO_2 and $\alpha_{Zr}(O)$ for uncoated and inner-coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst thermal ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C – Comparison with previous results obtained after one-sided steam oxidation of uncoated low-tin Zircaloy-4 [60] and using the same experimental methodology on uncoated M5_{Framatome} [47]

(a) Uncoated

(b) Inner-coated

Figure 23 - PQ hydrogen contents measured by neutron tomography (3D-maps and results plotted every 10° over the clad circumference) and by post-mortem IGFTC analysis of HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst thermal ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C followed by direct water quenching down to RT

(a) uncoated

(b) inner coated

Figure 24 - Through-thickness micro-hardness profiles measured at different axial locations of uncoated and inner coated HSn-HO-Zirc-4 clad segments that have experienced internal pressure ballooning and burst thermal ramp tests, and then subsequent HT steam oxidation for 600s at 1200°C + direct water quenching down to RT

(a)

(b)

(*) F/rod: failure load measured upon PQ tensile test extrapolated to full clad section [82]

Figure 25 - Relationships between maximum loads derived from PQ RCTs at 135°C and PQ tensile data from IRSN obtained on pre-hydrided Zirc-4 [82]), and:

(a) circumferentially averaged measured (interpolated) hydrogen concentrations

(b) interpolated prior- β_z micro-hardness and corresponding oxygen contents derived from equation 2

Figure 26 - Calculated PQ maximum tensile strength of prior- β_{Zr} at RT, using Le Hong et al.' correlation [86] for low-tin Zirc-4 vs. oxygen (*) and hydrogen (**) respective contents, plotted as a function of the maximum load derived from PQ RCTs carried out at RT or at 135°C on uncoated/coated HSn-HO-Zirc-4
 (*) derived from prior- β_{Zr} HV(0.01) measurements
 (**) interpolated (averaged) values from IGFTC measurements