



Bonded and additively manufactured crack retarders: A comparative study of damage tolerance properties

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ABSTRACT

Fatigue crack grows faster in the integral structure than in the riveted structure. This study compares the bonded, cold-sprayed and ultrasonically consolidated crack retarders to increase the fatigue life of integral airframe structure. The 2024-T351 aluminum M(T) specimen reinforced with crack retarders made of AISI 301 and 316L steels, Ti-6Al-4V titanium alloy and unidirectional CFRP laminate are evaluated in terms of fatigue life, delamination resistance and weight efficiency. Specimens were subjected to constant amplitude loading up to the final failure. A substantial decrease in fatigue crack growth rate was achieved using the ultrasonically consolidated AISI 301 crack retarder.

1. Introduction

The integral airframe structure saves weight and contains fewer stress concentrations but does not contain physical barriers against fatigue crack growth. The physical barrier is classified as a place in a structure where one part of a structure is covered by another part. If the physical barriers are non-existent, the fatigue crack can grow from one integral element to another without delay. It results in earlier fatigue failure of integral structure [1].

One of the techniques to slow down crack propagation is the application of so-called “crack retarders”. These selective reinforcements are usually made of materials with different mechanical properties than the base structure material. Their role is to reduce the crack opening through the bridging effect [2]. The crack retarder also acts as a stiffener and transfers the load from the substrate before the crack reaches the stiffener's position.

However, the crack retarder may start delaminating in front of the progressing crack tip, where the maximum tensile stresses are, and/or at the free ends, where the maximum shear stresses are. The retarder can still carry the load but is less effective due to the lack of shear transfer capability [3]. Crack retarders are located on one side of the airframe structure; the structure's central axis is shifted. As the unsymmetrical structure is tension loaded, the bending moments

emerge and cause out-of-plane deformation [4], which results in a stress increase at the crack tip. Suppose coefficients of thermal expansion of substrate and crack retarder deviate. In that case, the change in ambient temperature (manufacturing process, service conditions) promotes residual thermal stresses [5] that may affect stress at the crack tip. Another issue to be addressed is galvanic corrosion when combining dissimilar materials and crack retarder fatigue sensitivity.

Previous studies demonstrated a promising increase in fatigue life when using crack retarders made of ARALL fiber metal laminate [6], Al₂O₃/Al metal matrix composite [7], CFRP laminates [3,8], Ti-6Al-4V alloy [9], GLARE fiber metal laminate [10,11], cold-sprayed 7075 alloy [12] and ultrasonically consolidated Al₂O₃/Al metal matrix composite [13]. Furthermore, crack retarders were extensively evaluated in terms of residual thermal stresses after curing [14–16]; their durability (impact damage, thermal cycling) [17,18] and optimal design parameters [19].

A solid-state joint can be achieved between dissimilar metals using cold spray and ultrasonic consolidation. Cold spray is the technology that allows depositing thick metallic and non-metallic coatings. In this process, compressed and heated gas is used to carry particles through the de Laval nozzle. As the mixture expands, the velocity of particles increases. The particles then impact the substrate and create a metallurgical bond with the substrate and/or other particles [20]. Ultrasonic

Abbreviations: BCR, Bonded Crack Retarder; BCRs, Bonded Crack Retarders; CF, Carbon Fiber; CSCR, Cold-Sprayed Crack Retarder; CSCRs, Cold-Sprayed Crack Retarders; CTE, Coefficient of Thermal Expansion; UCCR, Ultrasonically Consolidated Crack Retarder; UD, Unidirectional

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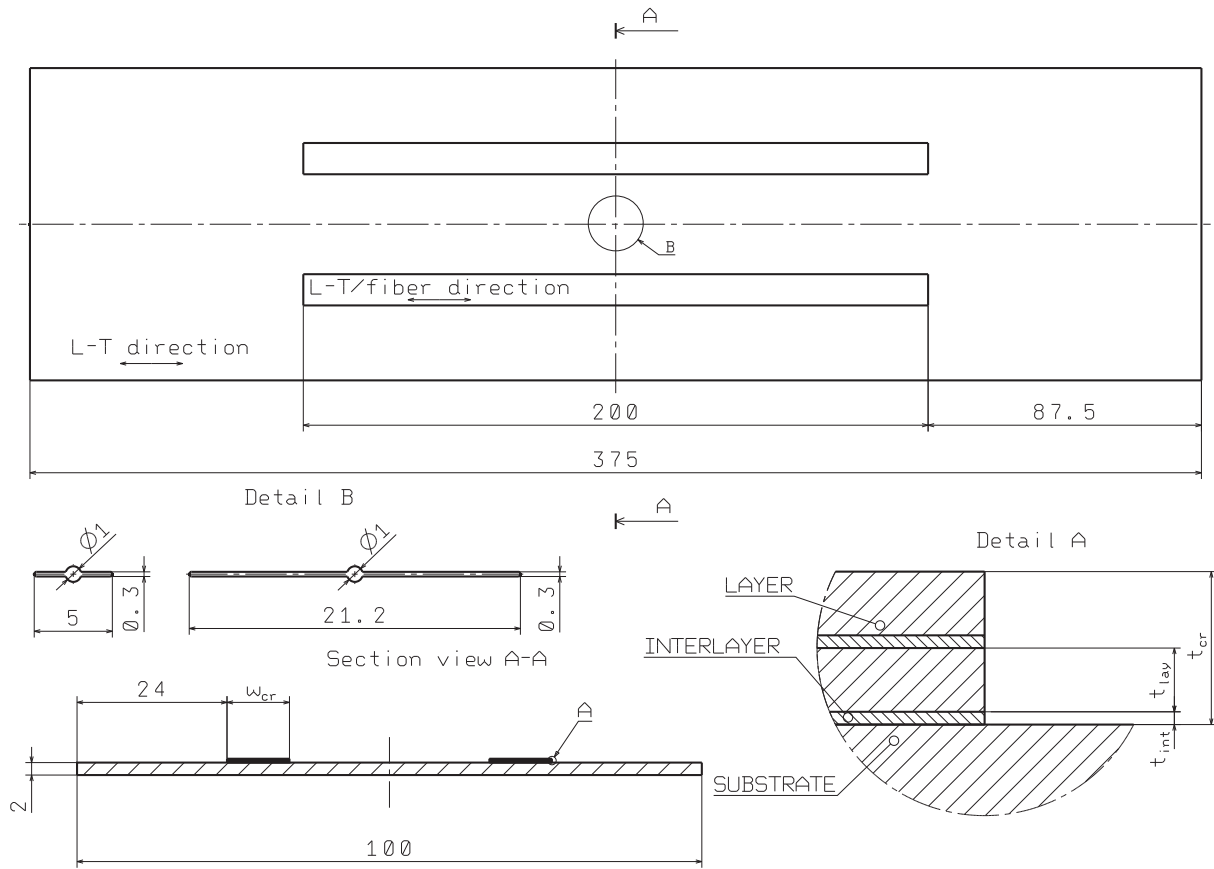


Fig. 1. Specimen shape. Dimensions in millimeters, not in scale. Specimens with CSCR and BCR: 5 mm long notch. Specimens with UCCR: 21.2 mm long notch.

Table 1
2024-T351 chemical composition [25].

Alloy	Al	Cr	Cu	Fe	Mg	Mn	Si	Ti	Zn
2024	Bal.	≤0.1	3.8	≤0.5	1.2	0.3	≤0.5	≤0.15	≤0.25
-T351			4.9		1.8	0.9			

consolidation is the process of continuous ultrasonic welding. It uses the mechanical vibrations and normal force of sonotrode to remove the oxide layer between two metallic foils. As the sonotrode rotates, it creates the metallurgical bond along the predefined path [21]. In as-built condition, the shear strength of ultrasonically consolidated Al-Ti joint [22, 23] and cold-sprayed AISI 316L-Al joint [24] has reached or exceeded the maximum shear strength of adhesive joints. We hypothesize that crack retarders manufactured using these two additive technologies might outperform bonded crack retarders. It has been demonstrated only for material combination (Al retarder on Al substrate) that due to the excellent adhesion of cold-sprayed crack retarders, the fatigue life of the specimen can be increased [12]. However, cold-sprayed crack retarders made of a different material from the substrate have not been evaluated in terms of their performance. This study also investigates the performance of ultrasonically consolidated crack retarders since only one study focused on the $\text{Al}_2\text{O}_3/\text{Al}$ metal matrix composite [13] has been found.

Due to the combination of the CTE close to the substrate's value and high elastic modulus, stainless steel was chosen as the material for the crack retarders. The combination of such material properties might increase the crack retarder's performance. For comparison purposes, the CF UD laminate was included in this study because it has proven outstanding crack-retarding abilities.

2. Manufacturing methods

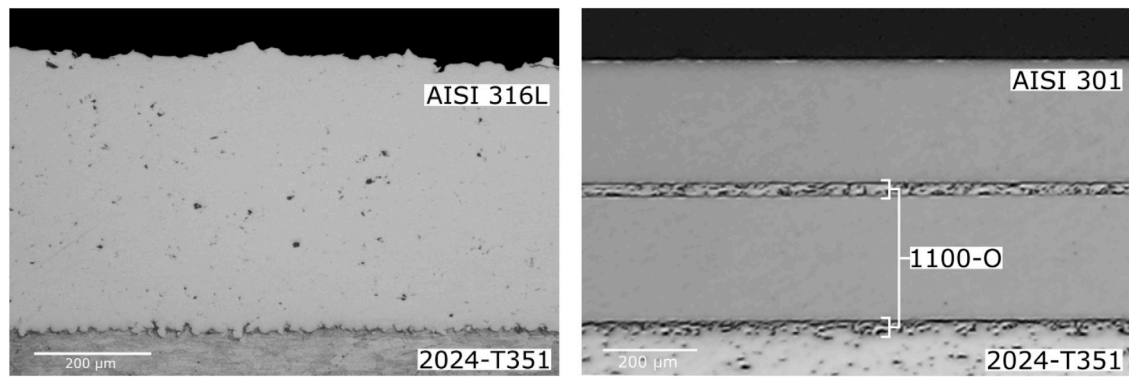
2.1. Substrate

The M(T) specimens (Fig. 1) were high-speed milled from the sheet made of 2024-T351 aluminum alloy (Table 1), which is widely used in various aerospace applications. The bare M(T) specimens were used as a substrate for bonded and cold-sprayed crack retarders and as a reference specimen for comparison purposes. The M(T) specimen width and length were chosen according to ASTM E647 [30] while the thickness of 2 mm represented the typical integral wing skin panel. The mechanical properties of substrate are shown in Table 2. In the case of specimens with ultrasonically consolidated crack retarders, the substrate was manufactured in the reverse order (as elaborated in Section 2.4). The geometry of substrate for consolidated crack retarders and the location of consolidated crack retarders are depicted in Fig. 1. Dimensions of each crack retarder and its constituents are presented in Table A.1.

2.2. Bonded crack retarders

2.2.1. Austenitic stainless steel

Previous studies on crack retarders for aerospace applications excluded steels due to the higher specific weight that could result in high fuel consumption and CO_2 emissions. However, the stainless steels possess the CTE close to the aluminum alloy in combination with excellent tensile strength and elastic modulus. Hence, the austenitic stainless steel AISI 301 in the work-hardened state was selected for this study (see Table 2, 3 for mechanical properties and chemical composition). The capability of retarding the crack growth was examined using specimens with one bonded steel layer and two bonded steel layers.



(a) 1-AISI316L-CSCR specimen: cold-sprayed AISI 316L crack retarder
(b) 2-AISI301-UCCR specimen: Ultrasonically consolidated AISI 301 crack retarder

Fig. 2. Micrograph of additively manufactured crack retarders.

Table 2

Mechanical and thermal properties of substrate, bonded crack retarder and adhesive.

Material		2024-T351 ^a	AISI 301	M10R/38% /UD150/CHS	Araldite® 2011
Young's modulus	E [GPa]	72.4	185 ^b	136 ^d /143 ^e	1.9 ^h
Density	ρ [g/cm ³]	2.77	7.88 ^c	1.57 ^f	1.05 ^h
CTE	α [10 ⁻⁶ /K]	23	17 ^c	1 ^g	–
Tensile strength	R_m [MPa]	470	1635 ^b	2168 ^d /2676 ^e	–
Yield strength	$R_{p0.2}$ [MPa]	325	1508 ^b	–	–
Elongation	A [%]	20	2.1 ^b	–	–

^a[25].

^b[26].

^c[27].

^dTwo layers. Tested according to ASTM D3039.

^eFive layers. Tested according to ASTM D3039.

^f[28].

^gTypical generic values.

^h[29].

Table 3

AISI 301 chemical composition [27].

Alloy	Fe	C	Mn	Si	P	S	Cr	Ni
AISI 301	Bal.	≤0.15	≤2	≤1	≤0.045	≤0.03	16 18	6 8

Table 4

SANDVIK 316L (15–38 μm) powder chemical composition [31].

Alloy	Fe	C	Cr	Ni	Mo	Si	Mn	S	P
AISI 316L	Bal.	≤0.03	16.0 18.0	10.0 14.0	2.0 3.0	≤1.0	≤2.0	≤0.03	≤0.045

Crack retarder surface pretreatment. The straps used for adhesive bonding of crack retarders were cut out of the sheet made of 0.255 mm thick AISI 301 steel and degreased with acetone. Subsequently, they were immersed in the solution containing 12.5% HF, 40.8% H₂O, and 46.7% HNO₃ for 20 min at the ambient temperature. In the end, rinsing with water and blow-drying with the 45 °C air took place.

Adhesive bonding. The crack retarders were bonded to the specimen surface using the two-component Araldite® 2011 structural adhesive. The specimen with one steel layer (1-AISI301-BCR) was cured for 24 h at the ambient temperature and 30 min at 80 °C. In the case of the specimen with two steel layers (2-AISI301-BCR), first, two pretreated straps were bonded together and cured. An excessive adhesive layer on the outer surfaces was removed. Then, the resulting BCRs were degreased with acetone and immersed in the solution containing H₂O, HF and HNO₃ for 20 min at the ambient temperature. Finally, bonding onto the substrate and curing were done. The same curing process was used as in the case of the 1-AISI301-BCR. Note that two specimens with two steel layers were manufactured; specimens 2-AISI301A-BCR and 2-AISI301B-BCR.

2.2.2. Unidirectional carbon fiber

To compare the performance of steel crack retarders with previously validated crack retarder material, the specimens with BCRs made of two (1-CFUD-BCR) and five layers (2-CFUD-BCR) of M10R/38%/UD150/CHS prepreg were prepared. The prepreg consists of the high-strength unidirectional 12 K carbon fiber fabric and the M10R epoxy resin matrix (see mechanical properties in Table 2).

Crack retarder manufacturing process. First, the prepreg roll was defrosted at the ambient temperature to avoid the condensation of air humidity. Then, the individual prepreg plies were cut out of the prepreg roll. After removing the release film from the prepreg plies, the desired number of prepreg plies was positioned in a single direction. Then the entire assembly was covered with the peel ply, bleeder cloth and vacuum bagging film. This assembly was autoclaved at 5 bar and 120 °C for 1 h. Finally, the cured sheets were cut with a hand saw to obtain 10 mm wide crack retarders.

Crack retarder surface pretreatment. Using peel ply, the crack retarder surfaces were modified to optimize the bond quality. Before bonding, they were degreased with acetone.

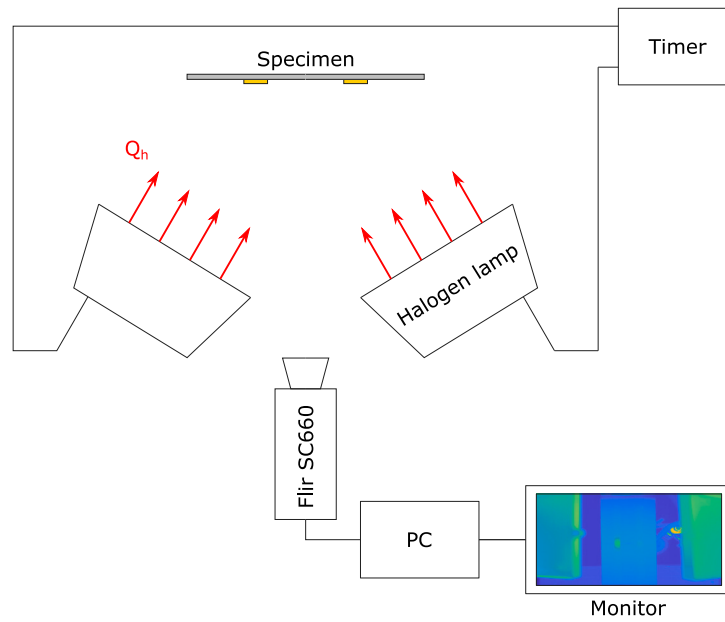


Fig. 3. Pulsed thermography apparatus.

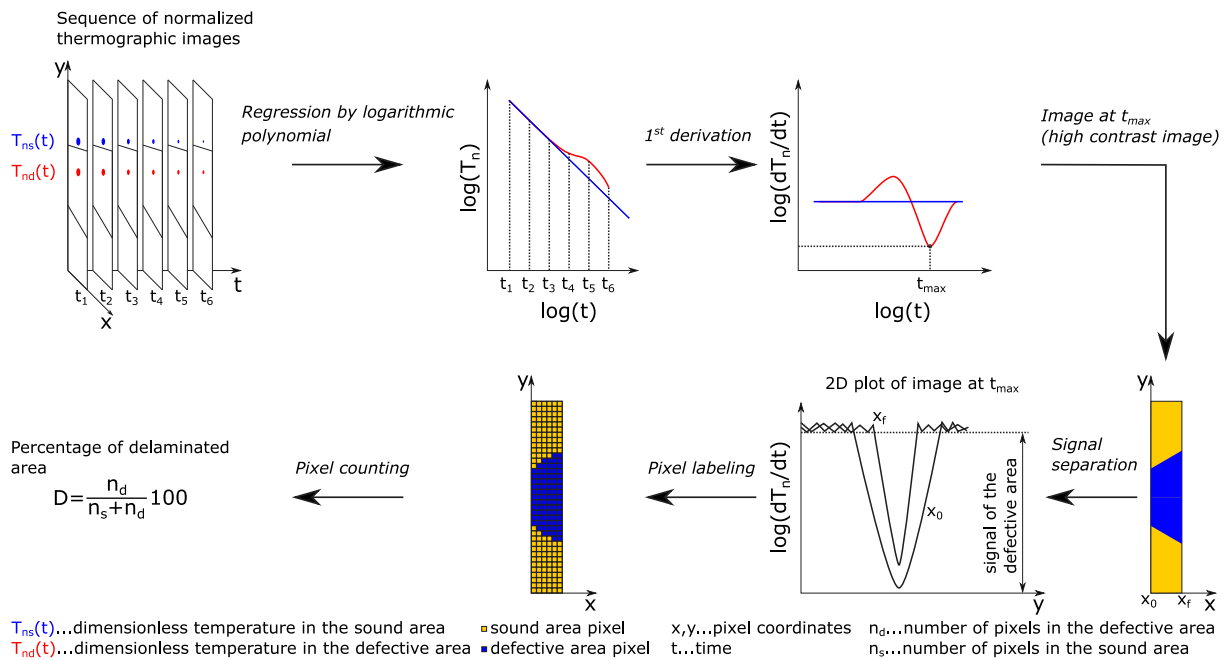


Fig. 4. Thermographic image processing.

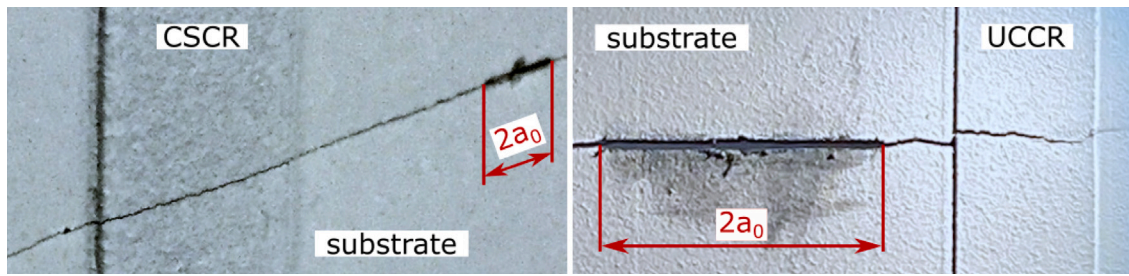
Table 5
 AP&C Ti-6Al-4V (15–45 μ m) powder chemical composition [32].

Alloy	Ti	Al	V	Fe	O	C	N	H	Y
Ti-6Al-4V	Bal.	5.50	3.50	0.05	0.14				
		6.75	4.50	0.25	0.16	≤ 0.02	≤ 0.02	≤ 0.010	< 0.005

Adhesive bonding. The CF UD crack retarders were bonded to the 2024-T351 substrate surface using the Araldite[®] 2011. The 1-CFUD-BCR specimen and 2-CFUD-BCR specimen were first cured for 24 h at the ambient temperature and then cured for 30 min at 80 °C. Admittedly, combining the carbon fiber laminate with aluminum alloy might lead to galvanic corrosion and the hot curing process introduced residual thermal stresses to the specimen.

2.3. Cold-sprayed crack retarders

To compare the bonded crack retarders with cold-sprayed ones, two specimens with alpha-beta alloy Ti-6Al-4V and two with austenitic stainless steel AISI 316L crack retarders were manufactured (see Tables 4 and 5 for powder properties). Crack retarders were deposited (Fig. 2(a)) using the Impact Spray System 5/11 and optimized process



(a) Specimen 1-AISI316L-CSCR. Cold-Sprayed Crack Retarder (CSCR). (b) Specimen 1-AISI301-UCCR. Ultrasonically Consolidated Crack Retarder (UCCR).

Fig. 5. Direct propagation of fatigue crack to crack retarder.

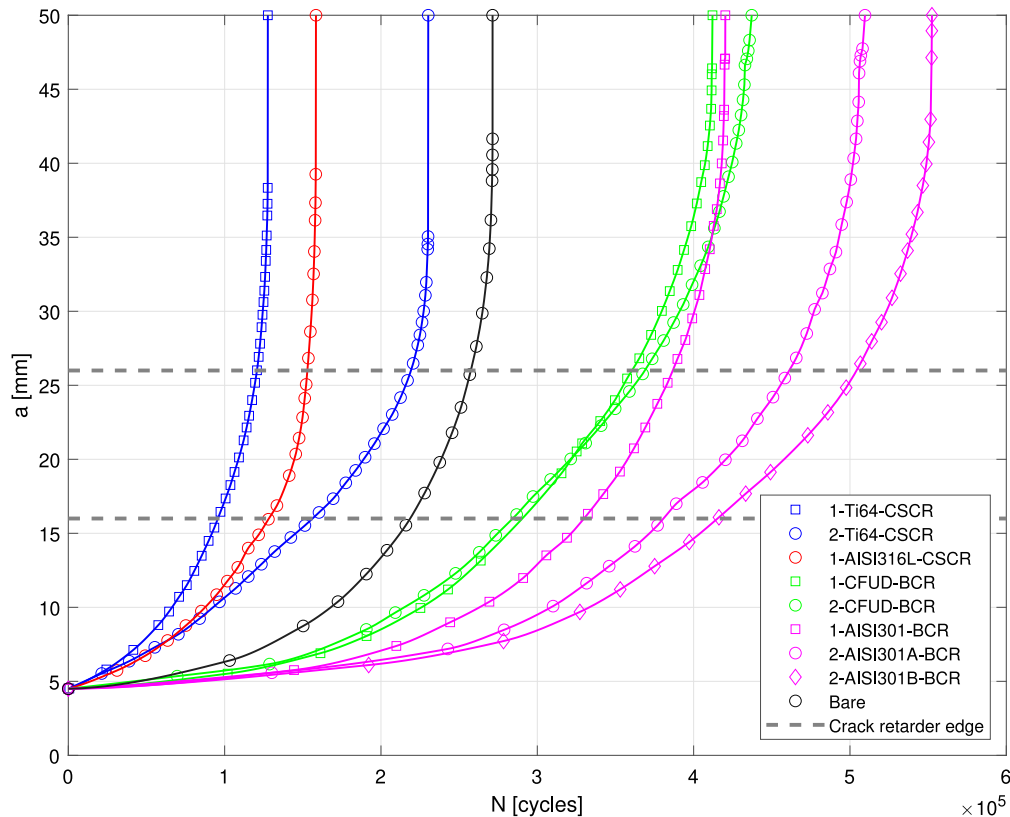


Fig. 6. Crack growth curves: bonded and cold-sprayed crack retarders; $\sigma_{max} = 60$ MPa, $R = 0.1$, $f = 15$ Hz.

parameters: stand-off distance 30 mm, step distance 1 mm, gun travel speed 0.5 m s^{-1} , gas (N_2) pressure 50 bar and temperature 1100°C . As overspray protection, the metal mask was utilized to obtain consistent width and length of crack retarders.

Cold-sprayed deposits in as-sprayed condition often possess lower mechanical properties (Table 6) than wrought counterparts. To increase retarders' load transfer capacity, their height was increased to 0.5 mm (1-AISI316L-CSCR and 1-Ti64-CSCR specimens) and 1.0 mm (2-AISI316L-CSCR and 2-Ti64-CSCR specimens).

2.4. Consolidated crack retarders

Two 2024-T351 M(T) specimens with ultrasonically consolidated crack retarders were manufactured. The capability of retarding the crack growth was examined for the specimens with crack retarders consisting of one and two steel layers. Crack retarders were made of AISI 301 steel; the same composition, thickness and batch as in the case of BCRs.

Table 6

Mechanical and thermal properties of cold-sprayed crack retarder.

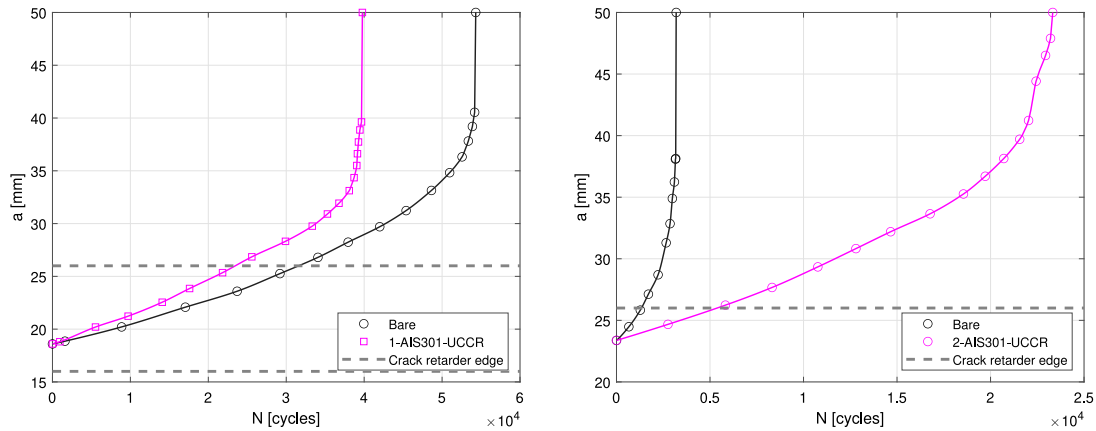
Material		Ti-6Al-4V	AISI 316L
Young's modulus	E [GPa]	69.7 ^a	182 ^a
Porosity	ϕ [%]	1.1/1.8 ^b	0.7/1.0 ^b
CTE	α [$10^{-6}/\text{K}$]	8.6 ^b	15.9 ^c
Tensile strength	R_m [MPa]	295 ^a	795 ^a
Yield strength	$R_{p0.2}$ [MPa]	—	765 ^a
Elongation	A [%]	0.05 ^a	0.05 ^a

^aASTM E8 Sheet-Type Subsize Specimens with 6×3 [mm] reduced section were used.

^bAt least five images of each cold-sprayed material were evaluated utilizing optical microscope and thresholding technique.

^c[27].

The substrate surface had to be machined before consolidation to be perfectly leveled and clean. For this reason, using the same batch of 2024-T351 bare M(T) specimens was not possible as in previous experiments. Before consolidation, the 6.35 mm thick 2024-T351 sheet was



(a) Specimen 1-AISI301-UCCR; $\sigma_{max} = 60$ MPa, $R = 0.1$, $f = 15$ Hz (b) Specimen 2-AISI301-UCCR; $\sigma_{max} = 100$ MPa, $R = 0.1$, $f = 15$ Hz.

Fig. 7. Crack growth curve: ultrasonically consolidated crack retarders.

Table 7

Mechanical and thermal properties of 1100-O interlayer used in consolidated crack retarder.

Material		1100-O
Young's modulus	E [GPa]	69.0 ^a
Density	ρ [g/cm ³]	2.71 ^a
CTE	α [10 ⁻⁶ /K]	23.6 ^a
Tensile strength	R_m [MPa]	90 ^a
Yield strength	$R_{p0.2}$ [MPa]	35 ^a
Elongation	A [%]	40 ^a

^a[33].

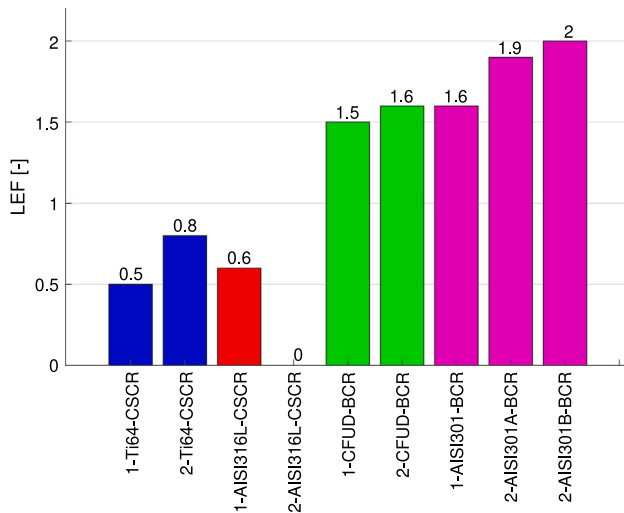


Fig. 8. Life extension factor: bonded and cold-sprayed crack retarders.

milled down to 6 mm. Crack retarders were consolidated on the milled surface. Specimens (1-AISI301-UCCR, 2-AISI301-UCCR) were then extracted from the sheet. Additionally, 2024-T351 bare M(T) specimens were extracted for comparison purposes and to avoid batch-to-batch inconsistency.

2.4.1. Austenitic stainless steel

There are two approaches to consolidate foils made of materials with a hard oxide film (e.g. stainless steels): increasing the process parameters (normal force, amplitude) and/or using the relatively soft interlayer. The former approach can cause the foil to bond with the

sonotrode or lead to the sonotrode's excessive wear [34]. The latter causes the interlayer material to flow around hard oxide film resulting in the bond between two materials with hard oxide film [35]. In this study, the 0.05 mm thick 1100-O interlayer was used between AISI 301 layers (Table 7). Crack retarders were deposited (Fig. 2(b)) using the Fabrisonic SL 7200, Ultrasonic Additive Manufacturing machine with the 9 kW ultrasonic welding system. The following optimized process parameters were used: 20 kHz frequency, 40 μ m amplitude, 21.2 mm/s longitudinal speed, 7500 N down force, anvil heated at 52 °C, titanium driver. The driver was placed between the sonotrode and the foil to be consolidated. It holds foil in place, transfers the horn's oscillations and prevents foil from bonding to the horn.

3. Test methods

Fatigue crack growth test. Prior testing, the initial notch ($2a_0 = 5$ mm) was machined to the specimen with bonded (CF UD, AISI 301), cold-sprayed crack retarders (Ti-6Al-4V, AISI 316L) and without crack retarder (Bare). Considerably longer notch ($2a_0 = 21.2$ mm) was machined to all specimens with ultrasonically consolidated crack retarders. The crack length was periodically measured on the specimen's unreinforced side using the traveling microscope with 32x magnification. The cyclic load was applied until the final failure of specimen. All specimens with bonded crack retarders, cold-sprayed crack retarders, without crack retarders and the specimen with ultrasonically consolidated crack retarders (1-AISI301-UCCR) were subjected to the crack propagation test with the following parameters: $\sigma_{max} = 60$ MPa, $R = 0.1$ and $f = 15$ Hz.

In the case of the 2-AISI301-UCCR specimen, the applied stress had to be increased. Even with the larger machined notch ($2a_0 = 21.2$ mm), the precracking in the 2-AISI301-UCCR specimen was unsuccessful at the $\sigma_{max} = 60$ MPa and $\sigma_{max} = 80$ MPa. The crack did not start growing after the 1.0e6 – 1.4e6 cycles. For this reason, the test was conducted at $\sigma_{max} = 100$ MPa, $R = 0.1$ and $f = 15$ Hz. At this stress level, the fatigue crack started to grow.

Delamination monitoring. Throughout the fatigue crack growth test, the thermographic inspection was carried out. After the optical excitation through 2x 400 W halogen lamps (Fig. 3), the delamination monitoring was performed by measuring the thermal response of the specimen using the Flir SC660. The thermal response was collected during heating and decay; while excitation took 5, 7 and 10 s. It appeared from the first processed thermal sequences that the highest contrast was acquired after 10 s excitation, so the analysis was performed only for this excitation time. Before testing, specimens were painted with white spray-paint ThermoSpray 500 to minimize the negative effect of reflective heat.

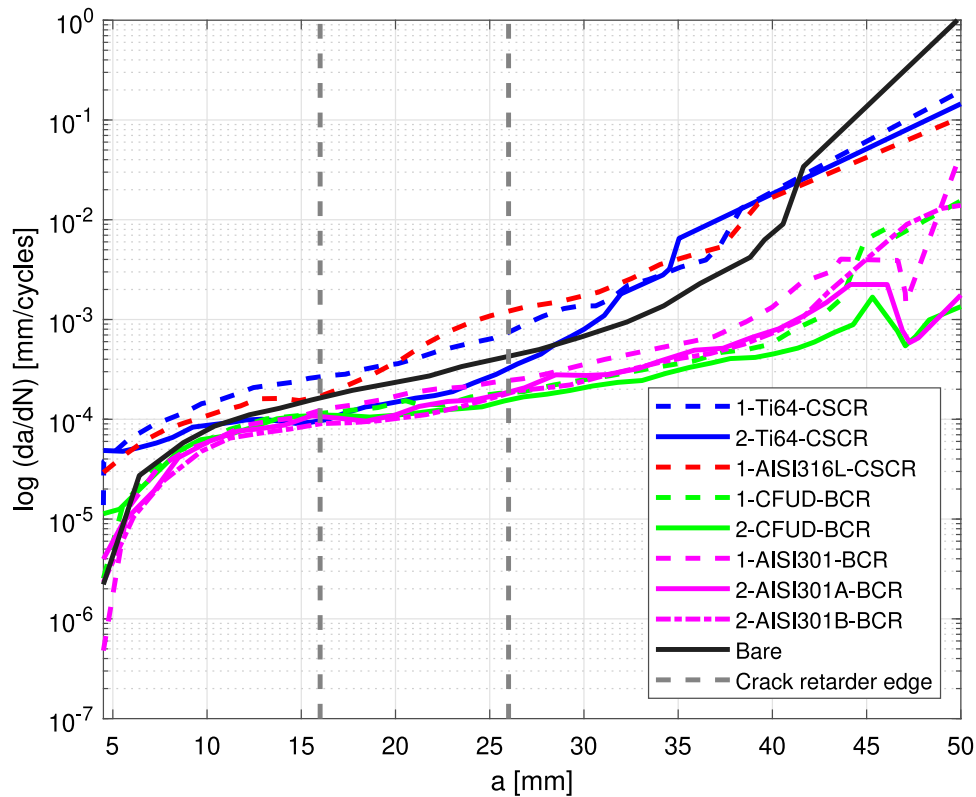


Fig. 9. Fatigue crack growth rate: bonded and cold-sprayed crack retarders.

Thermographic image processing. First, the decay sequence was extracted from the specimen's thermal response and normalized by dividing the thermal decay images by the image at the end of heating. Normalization reduced the effect of non-uniform heating and optical properties variability across the measured surface. The normalized thermal sequence was fitted with polynomials (Fig. 4) to reduce noise. By solving the first derivative of each polynomial, the contrast between the defective and the sound area was increased (i.e. Thermographic Signal Reconstruction method). From the first derivative, the time of high contrast image was found. The 2D plot was created from that image to find the defective area signal (sharp peak in the 2D plot). The pixels belonging to the defective and sound areas were sorted and counted. Finally, the delaminated area was calculated for each crack retarder.

4. Experimental results and discussion

Crack growth. During the crack propagation test, no crack initiation was observed in the BCRs. They broke¹ or disbond² shortly after the substrate failure. In cases of all cold-sprayed crack retarders, the crack propagated directly from the substrate to the crack retarders (Fig. 5(a)). In the case of ultrasonically consolidated crack retarders, the failure was contradictory. The fatigue crack in the specimen with one steel layer (1-AISI301-UCCR, $\sigma_{max} = 60$ MPa) directly propagated to the crack retarder (Fig. 5(b)). However, the crack in the specimen with two steel layers (2-AISI301-UCCR, $\sigma_{max} = 100$ MPa) remained in the substrate and the crack retarders broke immediately after the substrate failure. Measured crack lengths were averaged values of both crack sides and plotted against the number of cycles; thus, the crack propagation curves were obtained (Fig. 6, 7).

Regarding specimens with crack growth life longer than specimen without crack retarders (Bare), no significant delay can be observed in the crack retarder area (16–26 mm). The positive crack retarding effect manifests along the entire crack path. Similarly, any significant change in the crack growth rate cannot be observed in specimens possessing shorter fatigue life.

Fatigue life. The fatigue life improvement can be expressed by the life extension factor:

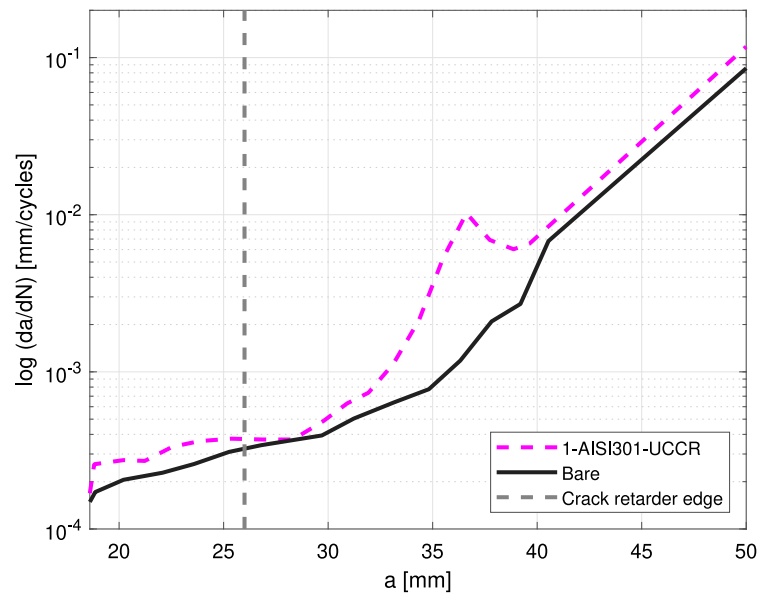
$$LEF = \frac{N_i - N_{bare}}{N_{bare}} \quad (1)$$

where N_i is the number of cycles to failure of specimen with crack retarders and N_{bare} is the number of cycles to failure of specimen without crack retarders. All bonded crack retarders resulted in a significant fatigue life extension. The greatest increase (1.9–2.0) was observed in the case of the specimen with two steel layers (Fig. 8). One can observe a relatively small difference between the fatigue crack growth curves of the specimen with two (1-CFUD-BCR) and five (2-CFUD-BCR) CF UD layers. It was probably caused by the evolution of residual thermal stresses during the curing process, which raised the mean stress and sped up the crack propagation in the specimen with thicker CF UD crack retarders. Regarding this observation, using more than two CF UD layers appears redundant. All cold-sprayed crack retarders dramatically decreased fatigue life. The most significant decrease, 50%, was observed in the fatigue life of the specimen with the Ti-6Al-4V crack retarders. The specimen with the thicker 316L crack retarder failed within the crack initiation period due to the rapidly growing crack at one of the ends of the crack retarder.

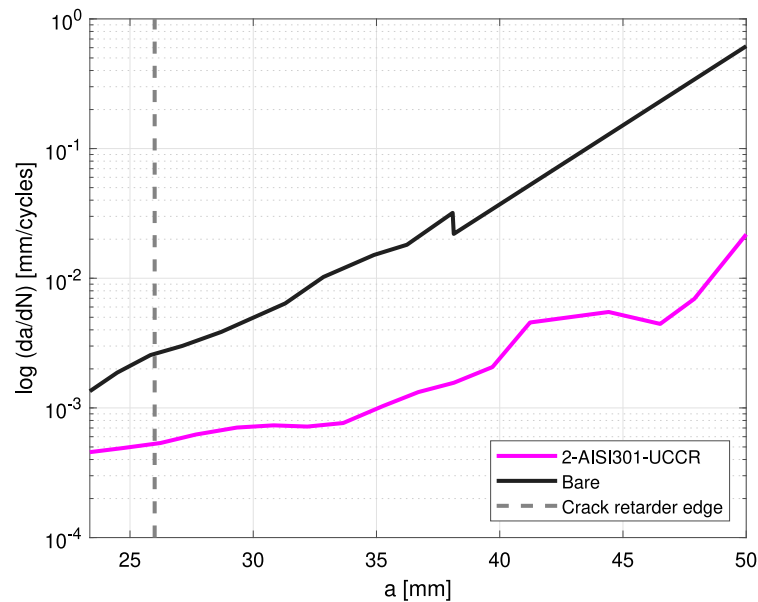
There are several explanations for the poor fatigue properties (i.e. decreased fatigue life and increased crack growth rate) of specimens with CSCRs. The substrate temperature increased due to its plastic deformation during cold spraying and heat transfer from the impinging jet. A combination of cooldown from such temperature and the crack retarder's CTE dissimilar from the substrate probably led to

¹ 1-AISI301-BCR, 1-CFUD-BCR

² 2-AISI301A-BCR, 2-AISI301B-BCR, 2-CFUD-BCR



(a) 1-AISI301-UCCR specimen.



(b) 2-AISI301-UCCR specimen.

Fig. 10. Fatigue crack growth rate: ultrasonically consolidated crack retarders.

the evolution of tensile residual stresses in the substrate. Compared with Liljedahl et al.'s study [5], similar out-of-plane deformation was observed in all specimens with CSCRs. When comparing the out-of-plane deformation in all specimens with CSCRs, the greatest deflection (20.4 mm) was measured in the center of the 2-Ti64-CSCR specimen. The temperature increase in the substrate certainly altered its mechanical properties. A noticeable drop in the substrate hardness was measured in all specimens with CSCRs. For example, a 22% decrease was recorded in the substrate of 1-AISI316L-CSCR specimen.

Specimen with one ultrasonically consolidated steel layer (1-AISI301-UCCR) failed earlier; its fatigue life was 30% lower than the specimen without crack retarder (Bare). However, the specimen with two steel layers (2-AISI301-UCCR) experienced the most significant fatigue life increase by a life extension factor of 7.3. Note that in this

specimen, the pre-crack length was five times greater and σ_{max} was 1.66 times greater compared to the specimens with BCRs and CSCRs.

Fatigue crack growth rate. Since the beginning of the test, the specimens with CF UD and steel crack retarders maintained the crack growth rate below the growth rate of the bare specimen (Fig. 9). Unfortunately, specimens with cold-sprayed crack retarders possessed a higher rate in the entire crack length range, resulting in shorter fatigue life. Although the crack retardation in the 2-Ti64-CSCR specimen can be identified in the (11–28) mm crack length range, the specimen failed earlier than the bare one because the crack grew significantly faster in other regions.

The fatigue crack growth rate in the specimen with one steel layer (1-AISI301-UCCR) was higher in the whole crack length range (Fig. 10(a)), which resulted in shorter fatigue life. On the contrary,

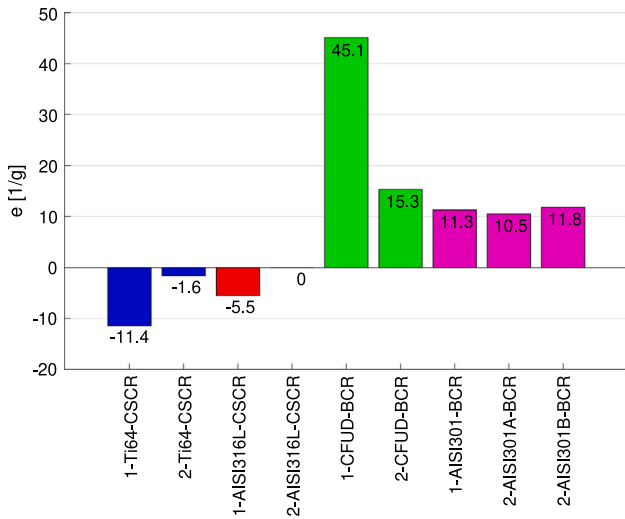


Fig. 11. e factor: bonded and cold-sprayed crack retarders.

the specimen with two steel layers possessed fatigue crack growth rate significantly lower compared to the specimen without crack retarders (Fig. 10(b)).

The percentage of life improvement with respect to the bare specimen per unit weight of crack retarder can be expressed by a following parameter:

$$e = \frac{N_f - N_{bare}}{N_{bare}} \frac{100}{m_{strap}} \quad (2)$$

where m_{strap} is the crack retarder weight. Obviously, the BCR made of two CF UD layers can significantly extend fatigue life while maintaining the relatively low weight of specimen (Fig. 11). One can also observe that the thinner BCRs were much more effective than the thicker ones. This behavior was already proved by Boscolo [19] and Molinari [36]. Specimens with steel BCRs did not perform in the same way. Compared to carbon fiber crack retarders, their total thickness was lower. They probably did not experience secondary bending to the extent that would have lowered performance. The greatest fatigue life increase with respect to the crack retarder unit weight, $82.2\% \text{ g}^{-1}$, was observed in the ultrasonically consolidated crack retarder with two steel layers (2-AISI301-UCCR).

Delamination propagation. Each specimen contains two crack retarders and potentially two delaminated areas. Thus, delaminated areas were averaged to obtain delamination growth curves. All bonded crack retarders started delaminating at the end of the specimen's fatigue life. The delamination in specimens with carbon fiber BCRs started later than in specimens with steel BCRs (Fig. 12). The steel crack retarders delaminated with a continually increasing delamination rate. The earlier onset of delamination in specimens with steel crack retarders can be attributed to weaker joint strength. The fracture surface between the CF UD crack retarder and the substrate showed adhesive-cohesive failure. On the contrary, the fracture surface between the steel crack retarder and the substrate was a purely adhesive failure. It suggests that if adhesive bond strength (AISI 301/substrate) is increased, the fatigue retardation effect may be increased.

Although several interface defects were observed in all specimens with CSCRs, the crack retarders did not delaminate. The crack grew directly to the crack retarder without premature delamination, probably due to the poor fatigue properties of the as-sprayed crack retarders. The poor fatigue properties of the cold-sprayed deposits were recently observed by Čížek et al. [37]. In their study, the pure metallic specimens (Ti, Ni, Al and Cu) possessed greater fatigue crack growth rates than their cold-rolled counterparts.

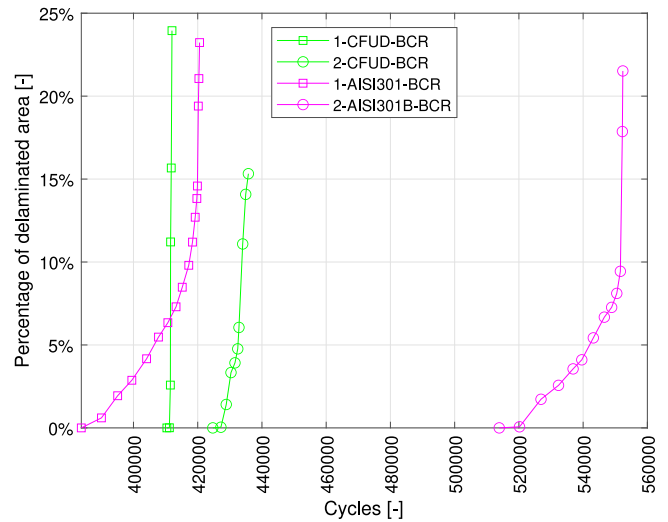


Fig. 12. Delamination growth: bonded crack retarders.

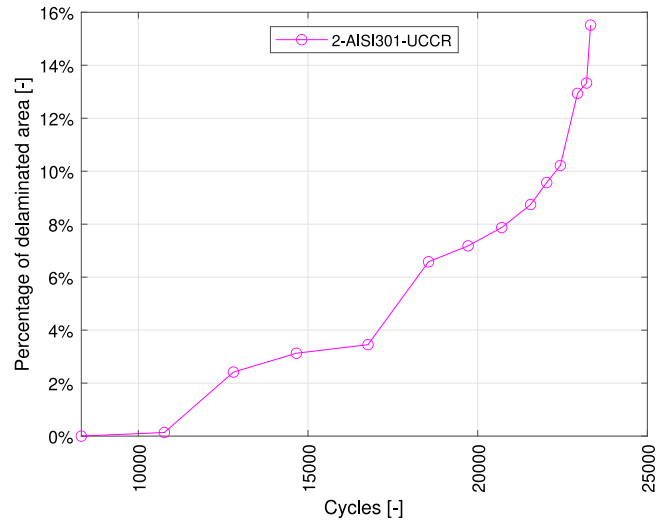


Fig. 13. Delamination growth: ultrasonically consolidated crack retarder.

Samely, the ultrasonically consolidated crack retarder (1-AISI301-UCCR) did not delaminate, the crack grew directly into him. Utilizing the ultrasonic consolidation, the joint stiffness was increased and more stress was transferred from the substrate to the crack retarder. Such stress levels allowed the crack to nucleate. The delamination in ultrasonically consolidated crack retarders (Fig. 13) followed a similar trend as in the bonded crack retarders (Fig. 12). Additionally, the delamination in the consolidated crack retarder possessed a triangular shape that is common in the bonded crack retarders³ [38].

5. Conclusions

This study investigated the crack retarding behavior and delamination progress of bonded, cold-sprayed and ultrasonically consolidated crack retarders. The cold-sprayed (Ti-6Al-4V, AISI 316L) and bonded crack retarders (AISI 301, CF UD) were tested at $\sigma_{max} = 60 \text{ MPa}$, $R = 0.1$ and $f = 15 \text{ Hz}$. The conclusions can be drawn:

³ Delamination growth in the left crack retarder; 2-AISI301-UCCR specimen and 2-AISI301B-BCR specimen.

1. Stainless steel bonded crack retarders performed better than carbon fiber ones in terms of life extension factor. The most significant increase in fatigue life was identified in specimens with two AISI 301 layers. The greatest percentual life increment with respect to the weight of crack retarder was identified in the specimen with two CF UD layers. However, residual thermal stresses were most likely induced in specimens with carbon fiber crack retarders because a hot curing process was used.
2. Cold-sprayed crack retarders made of 316L and Ti-6Al-4V could not slow down crack propagation. Cold-sprayed deposit formed an integral part with the substrate unlike the adhesively bonded crack retarder. Therefore, the fatigue crack grew directly from the substrate to the cold-sprayed crack retarder. After manufacturing, the most significant out-of-plane deformation was identified in specimens with cold-sprayed crack retarders. Fatigue crack acceleration might be owing to the residual thermal stresses. Because cold-sprayed crack retarders decreased fatigue life and increased the crack growth rate in the substrate, the cold spray technology is not suitable for depositing crack retarders.

Conclusions regarding ultrasonically consolidated crack retarders consisted of one steel layer (AISI 301) tested at $\sigma_{max} = 60$ MPa, $R = 0.1$, $f = 15$ Hz and two steel layers tested at $\sigma_{max} = 100$ MPa, $R = 0.1$ and $f = 15$ Hz are subsequently enumerated:

1. Ultrasonically consolidated crack retarder with two steel layers demonstrated the greatest crack retarding effect. A life extension factor of 7.3 was achieved; this corresponded to the life increase with respect to the weight of crack retarder of $82.2\% g^{-1}$. However, the crack retarder with a single steel layer performed worse than the specimen without crack retarder.
2. Pulsed thermography with thermographic signal reconstruction proved to be the relevant approach to measure delamination extent in ultrasonically consolidated crack retarders. The delamination growth in ultrasonically consolidated crack retarders followed the same trend as in bonded ones. Additionally, triangular delamination was recognized between the substrate and ultrasonically consolidated crack retarder.

Previous studies on crack retarders excluded steels due to the higher specific weight that might result in high fuel consumption and CO_2

emissions of an aircraft. This study filled a knowledge gap on the performance of stainless steel crack retarders manufactured using three distinctive technologies. However, the empirical results reported herein are subject to several limitations. Residual stresses were not measured in any specimen. Bonded and ultrasonically consolidated crack retarders were not examined for potential manufacturing defects. The residual stresses and defects might have a substantial effect on the results. Moreover, galvanic corrosion might be an issue when combining aluminum alloy and carbon fiber crack retarder.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

The processed thermographic data required to reproduce these findings are available to download from Mendeley Data: <http://dx.doi.org/10.17632/xvmwtj8hr9.1>.

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Appendix

See Table A.1.

Table A.1
Test matrix.

Specimen name	Crack retarder constituents					Crack retarder				Substrate		Loading parameters	
	No. of layers	Layer	Thickness (t_{lay}) [mm]	No. of interlayers	Interlayer	Length [mm]	Width (w_{cr}) [mm]	Thickness (t_{cr}) [mm]	Manufacturing temperature [$^{\circ}C$]	Material and geometry	Dimensions [mm]	Maximum stress (σ_{max}) [MPa]	Cyclic loading
Bare	0	–	0	0	–	0	0	0	RT	2024-T351 M(T)	$375 \times 100 \times 2$	60	$R = 0.1$ $f = 15$ Hz RT
1-AISI301-BCR	1	AISI 301	0.26	1	Araldite® 2011	200	10	0.44	80			60	
2-AISI301A-BCR	2	AISI 301	0.26	2	Araldite® 2011	200	10	0.60	80			60	
2-AISI301B-BCR	2	AISI 301	0.26	2	Araldite® 2011	200	10	0.55	80			60	
1-CFUD-BCR	2	M10R/38%/UD150/CHS	0.16	1	Araldite® 2011	200	10.2	0.53	80			60	
2-CFUD-BCR	5	M10R/38%/UD150/CHS	0.16	1	Araldite® 2011	200	10.2	1.08	80			60	
1-AISI316L-CSCR	3	AISI 316L	0.20	0	–	200	11.1	0.48	–			60	
2-AISI316L-CSCR	6	AISI 316L	0.20	0	–	200	11.2	0.96	–			60	
1-Ti64-CSCR	3	Ti-6Al-4V	0.20	0	–	200	11.2	0.51	–			60	
2-Ti64-CSCR	6	Ti-6Al-4V	0.20	0	–	200	11.3	1.02	–			60	
Bare	0	–	0	0	–	0	0	0	RT			60	
1-AISI301-UCCR	1	AISI 301	0.26	1	1100-O	200	10.2	0.29	52			60	
Bare	0	–	0	0	–	0	0	0	RT			100	
2-AISI301-UCCR	2	AISI 301	0.26	2	1100-O	200	10.2	0.71	52			100	

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