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SUSTAINED-LOAD CRACKING OF TITANIUM: A SURVEY
OF 6A1-4V ALLOYS

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Sustained-Load Cracking of Titanium

A Survey of 6Al-4V Alloys

G. R. YODER, C. A. GRIFFIS, AND
T. W. CROOKER

*Strength of Metals Branch
Metallurgy Division*

August 24, 1973



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ABSTRACT

This survey reveals that degradation in load-carrying capacity owing to sustained-load cracking (SLC) is widespread and serious in alloys of the Ti-6Al-4V family. Each of eight different 1-in. (2.5-cm) thick plates tested in air at room temperature exhibited the effect, with degradations ranging up to 35%. The amount of degradation does not seem to correlate with differences in hydrogen content, processing variables, or strength and toughness levels; however, the susceptibility to SLC is orientation dependent. Invalidity of several of the rising-load fracture toughness values, with respect to plane-strain criteria, makes quantitative comparison of degradations uncertain. However, the threshold stress-intensity factor below which SLC failures will not occur, K_{It} , clearly represents a more conservative fracture-safe design parameter than the plane-strain fracture toughness K_{Ic} , which cannot account for insidious time-delayed failures owing to SLC.

PROBLEM STATUS

This report completes one phase of the problem; work on other aspects of the problem is continuing.

AUTHORIZATION

NRL Problems M01-25 and M01-28
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NOMENCLATURE

A	lumped material constant
a	crack depth
B	plate thickness
C	constant (4.12 for cantilever loading, 3.80 for three-point bending)
c	concentration of hydrogen at the crack tip
c₀	concentration of hydrogen in the bulk
2c	length (major axis of surface crack)
D	specimen height; diffusivity
D₀	diffusion constant
\dot{K}	time derivative of stress-intensity factor
k	Boltzmann's constant
K_{Ic}	plane-strain fracture toughness
K_{Ii}	stress-intensity factor, based on initial crack length
K_{Ir}	stress-intensity factor, fracture resistance from rising-load test
K_{It}	threshold stress-intensity factor below which SLC failures will not occur
K_Q	stress-intensity factor, defined by ASTM-E399 5% secant offset method, from load displacement record
L	constant with dimension of length
M	applied bending moment
\dot{P}	time derivative of load
P_{max}	maximum load
P_Q	load defined by ASTM-E399 5% secant offset method, from load displacement record

PTC	part-through surface-cracked tension specimen
ppm	parts per million by weight
Q	flaw shape factor
r	radial polar coordinate
S	short transverse direction (thickness)
SEN	single-edge-notch bend specimen
SLC	sustained-load cracking
T	absolute temperature; long transverse direction
t_f	time to failure
\bar{v}	drift velocity
V	interaction energy between diffusing hydrogen and the crack-tip stress field
α	$1 - a/D$
Δ	SLC degradation in load-carrying capacity, in percent
θ	angular polar coordinate
σ	gross section stress
σ_{ys}	0.2% offset yield strength
τ	incubation time
∇	gradient

SUSTAINED-LOAD CRACKING OF TITANIUM: A SURVEY OF 6A1-4V ALLOYS

INTRODUCTION

While titanium alloys may be attractive for naval structural applications on the basis of high strength-to-density ratio, among other features, many exhibit potentially catastrophic crack (or flaw) extension while under static loads. Although such crack growth is usually attributable to stress-corrosion cracking, where aggressive environments are involved, it has also been reported for inert environments. In the latter case, crack growth has been referred to as sustained-load cracking (SLC), and has in several cases been attributed to the presence of interstitial hydrogen. The present effort to investigate SLC was motivated by the implications of this form of cracking with respect to structural life and proof testing. Results from the initial phase of this work, concerned primarily with determination of the SLC effect, are contained herein together with preliminary ramifications as to mechanism. Alloys examined, nominally from the 6A1-4V system, include both commercially and specially processed plates of widely divergent processing histories and interstitial contents.

The SLC phenomenon can be described with fracture mechanics parameters in the same manner that crack extension under static loading in an aggressive environment has been characterized for stress-corrosion cracking (1-3). Thus, if a material susceptible to SLC is loaded to initial stress-intensity K_{II} levels above a threshold level K_{IIc} , it will fail after a delay time t_f which decreases with increasing K_{II} . Delay times associated with this form of subcritical crack growth sometimes amount to several hours, and the threshold levels are as much as 35% less than the fracture resistance implied from standardized fracture toughness tests of only a few minutes' duration.

Awareness of SLC in titanium alloys dates back to the mid-1950's, when Burte *et al.* (4) demonstrated the time-delayed failure phenomenon with room-temperature stress-rupture tests of notched specimens in ambient air. The results of their work strongly implied that SLC behavior is due to hydrogen-assisted cracking, inasmuch as the phenomenon disappeared when the hydrogen content was reduced below a critical level known as the hydrogen tolerance. Later experiments by Sandoz (5) and Meyn (6), performed with fatigue-precracked specimens, suggested that the tolerance levels in titanium alloys based on the notched specimens were grossly overestimated, owing to the notch acuity effect demonstrated by Troiano (7), who found that the sharper the notch of the test specimen, the lower the apparent hydrogen content required for SLC of steels. Other reports of SLC in titanium alloys in ambient air have been made by Chu (8); Lane, Cavallaro, and Morton (9); and others (10).

PROCEDURE

Two specimen types were employed, the part-through surface cracked (PTC) tension specimen and the single-edge-notch (SEN) bend specimen, illustrated in Fig. 1. The latter

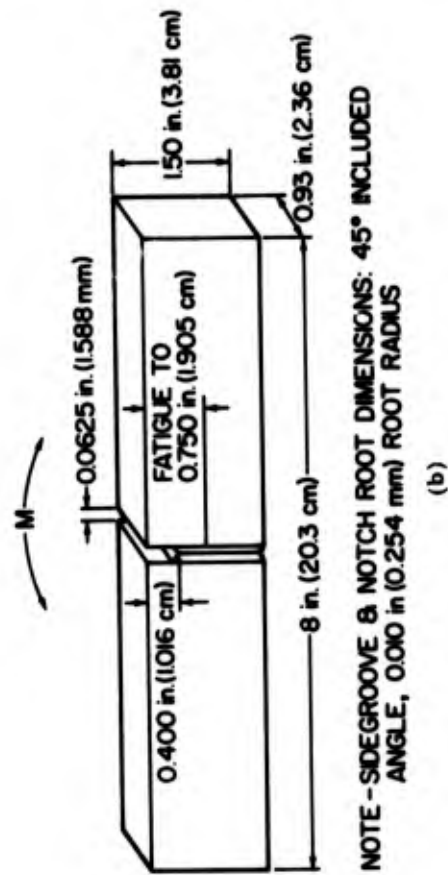
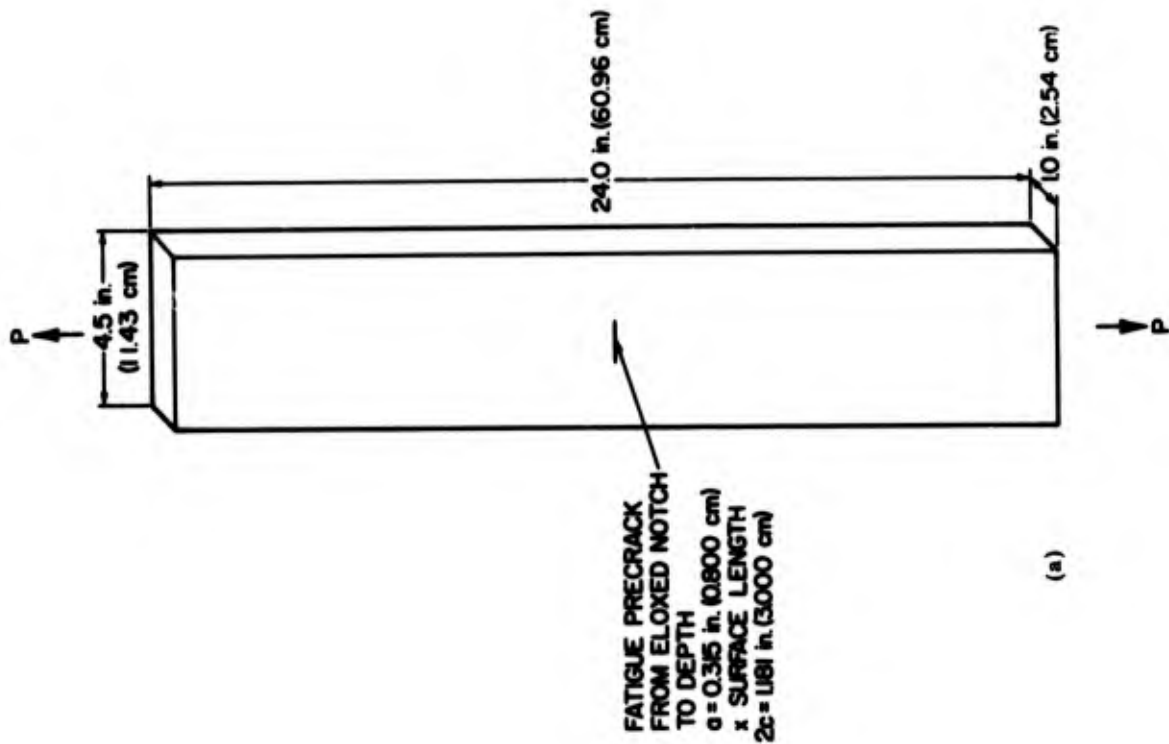


Fig. 1—Specimen geometries: (a) part-through cracked (PTC) tension specimen,
(b) single-edge-notch (SEN) bend specimen

type was used for the rising-load three-point bend test. For the SEN bend specimens, the K_{II} were computed from

$$K_{II} = CM \frac{\sqrt{\frac{1}{\alpha^3} - \alpha^3}}{BD^{3/2}} \quad (1)$$

where

$C = 4.12$ for cantilever loading (11) and 3.80 for rising-load bend tests (12)

$M =$ applied bending moment

$B =$ specimen thickness

$D =$ specimen height

$\alpha = 1 - a/D$ ($a =$ initial crack length).

For the PTC tension specimens (13),

$$K_{II} = 1.1 \sigma \sqrt{\pi \frac{a}{Q}} \quad (2)$$

where

$\sigma =$ gross section stress

$Q =$ flaw shape factor.

While all specimens were cut from 1-in. (2.5-cm) thick plate, different crack orientations were used for the two specimen types, viz. TS and TL (14) for the PTC and SEN specimens, respectively.

The SEN specimens were sidegrooved to a depth of $0.05B$ per side; thus while B (gross) = 0.93 in. (2.36 cm), B (net) = 0.84 in. (2.13 cm) was used for calculations with Eq. (1). Other SEN specimen dimensions were $D = 1.50$ in. (3.81 cm), $a \approx 0.750$ in. (1.91 cm), with specimen length = 8 in. (20.3 cm). For three-point bend tests, a span of 6 in. (15.24 cm) between rollers was used, while a loading rate of $\dot{P} \approx 8.6$ kips/min (38.2 kN/min) [$\dot{K} \approx 90$ ksi $\sqrt{\text{in.}}/\text{min}$ (100 MPa $\cdot\text{m}^{1/2}/\text{min}$)] was employed. The SEN specimens were fatigue precracked and equipped with displacement gages spanning the crack notches, in accord with the ASTM specifications (15). Step loading was not employed in any of the dead-weight cantilever tests for reasons described by Chu (8); rather, the full load was applied in a single application. Furthermore, a few specimens which displayed crack arrest were retested only after refatiguing the blunted crack tip to a point safely beyond the bounds of the plastic zone.

The PTC specimens were 24 in. (60.96 cm) long, 1 in. (2.54 cm) thick, and either 4.0 in. (10.16 cm) wide for rising-load tests or 4.5 in. (11.43 cm) for sustained-load tensile tests. Unless otherwise indicated, these specimens were loaded at the rate $\dot{P} = 500$ kips/s to either an appropriate level of sustained load or to failure in a rising-load test. These specimens were initially notched by electrical discharge machining, and then fatigue precracked to give "thumbnail" crack dimensions of approximately $a/2c = 0.315$ in. (0.800 cm)/1.181 in. (3.000 cm), where a is the crack length in the T direction of the plate and $2c$ is the major axis of the crack in the S direction. Fatigue precracking was accomplished by cycling the specimen between 0 and 225 kips (1000 kN) tension.

RESULTS

All plates tested exhibited SLC degradation in load-carrying capacity, more than 30% in some cases. In all instances, the greater portion of the apparent degradation occurred within a few hours of testing under sustained-load conditions. Results are summarized in Figs. 2-9 and Table 1. The table shows that materials examined ranged in (a) hydrogen contents from 35 to 80 parts per million by weight (ppm), (b) oxygen contents from 500 to 1600 ppm, (c) yield strengths (σ_{ys}) from 115 ksi (802 MPa) to 144 ksi (1005 MPa), (d) rolling temperatures from 1400°F (1033°K) to 1900°F (1311°K), and (e) forging temperature ranges from $\alpha + \beta$ to β . Each SLC curve in Figs. 2-9 is comprised of data points from both rising-load and sustained-load tests, the former of which are denoted by solid points, the latter with hollow points. From each curve the percentage of SLC degradation is measured as

$$\Delta = \left(1 - \frac{K_{It}}{K_{Ir}} \right) \times 100, \quad (3)$$

where K_{Ir} is the value of K_{II} from the rising-load test.

These degradations have been recorded in Table 1 together with other pertinent parameters which relate to the level of constraint represented by these tests. It is noted in the table that there are two possible values of K_{Ir} to choose from, viz. K_Q and $K_{I\max}$. The former is determined by the standard ASTM method from load-crack opening displacement traces (15), whereas the latter is calculated by using the maximum load value P_{\max} from the same trace. As shown in the table, however, in most cases K_Q fails to meet the criteria for valid K_{Ic} numbers (14), viz.,

$$\frac{P_{\max}}{P_Q} \leq 1.10 \quad (4)$$

$$2.5 \left(\frac{K_Q}{\sigma_{ys}} \right)^2 \leq B, a, D - a, \quad (5)$$

and is therefore a dubious parameter. Moreover, such an invalid K_Q number is not consistent with data from the sustained-load, cantilever bend tests. For example, in the case of alloy D4, $K_Q \approx 68 \text{ ksi}\sqrt{\text{in.}}$ (75.6 MPa-m^{1/2}), while $K_{It} \approx 70 \text{ ksi}\sqrt{\text{in.}}$ (77.9 MPa-m^{1/2}), and breaks appear at $K_{II} = 73$ and 77 ksi√in. (81.2 and 85.7 MPa-m^{1/2}) in the sustained-load cantilever bend results, as displayed in Fig. 7. On the other hand, $K_{I\max} = 80 \text{ ksi}\sqrt{\text{in.}}$ (90.0 MPa-m^{1/2}) is consistent with the cantilever bend data. To further

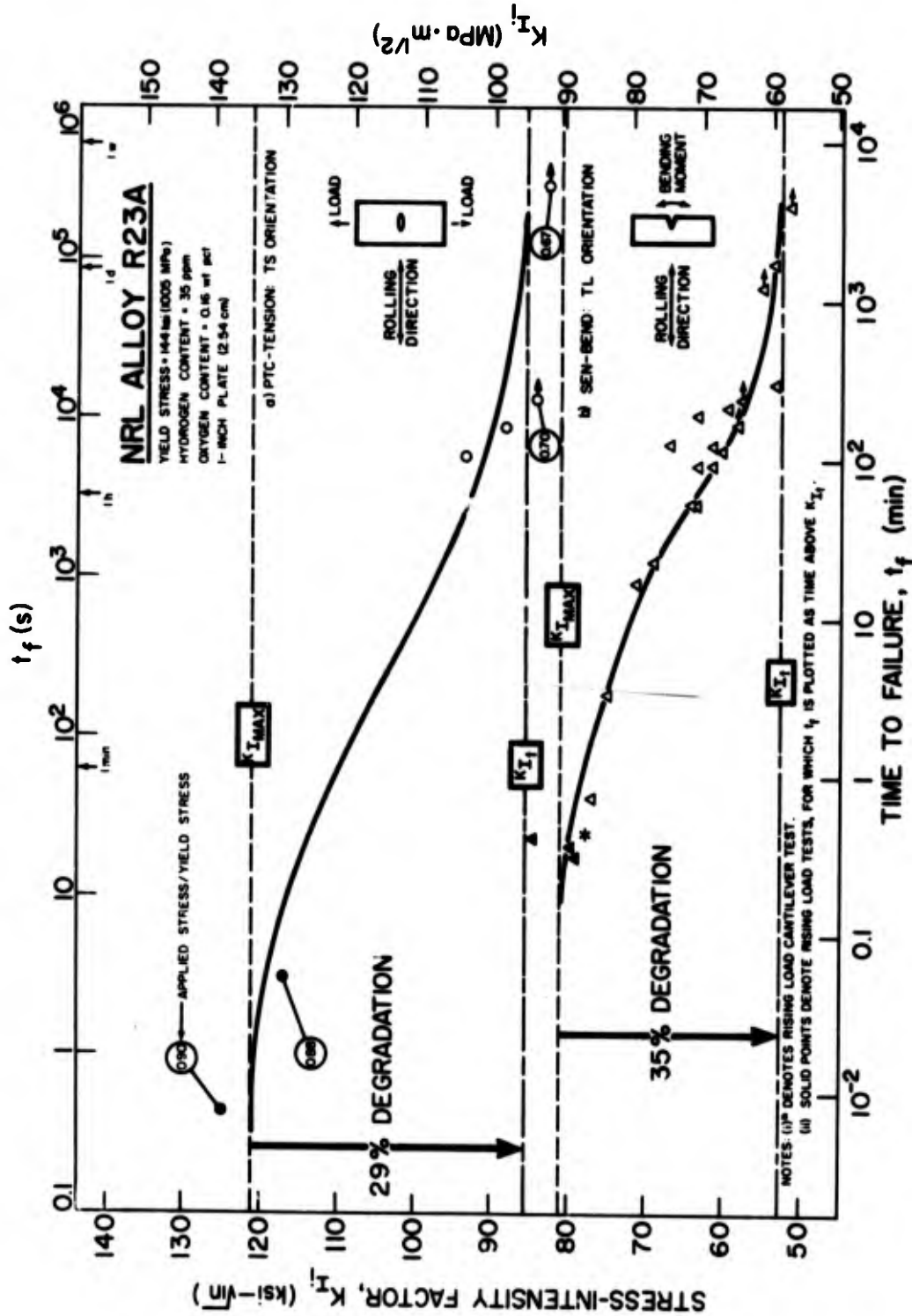


Fig. 2—Sustained load cracking in NRL alloy R23A in ambient, room-temperature air:
 (a) PTC tension specimens, (b) SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

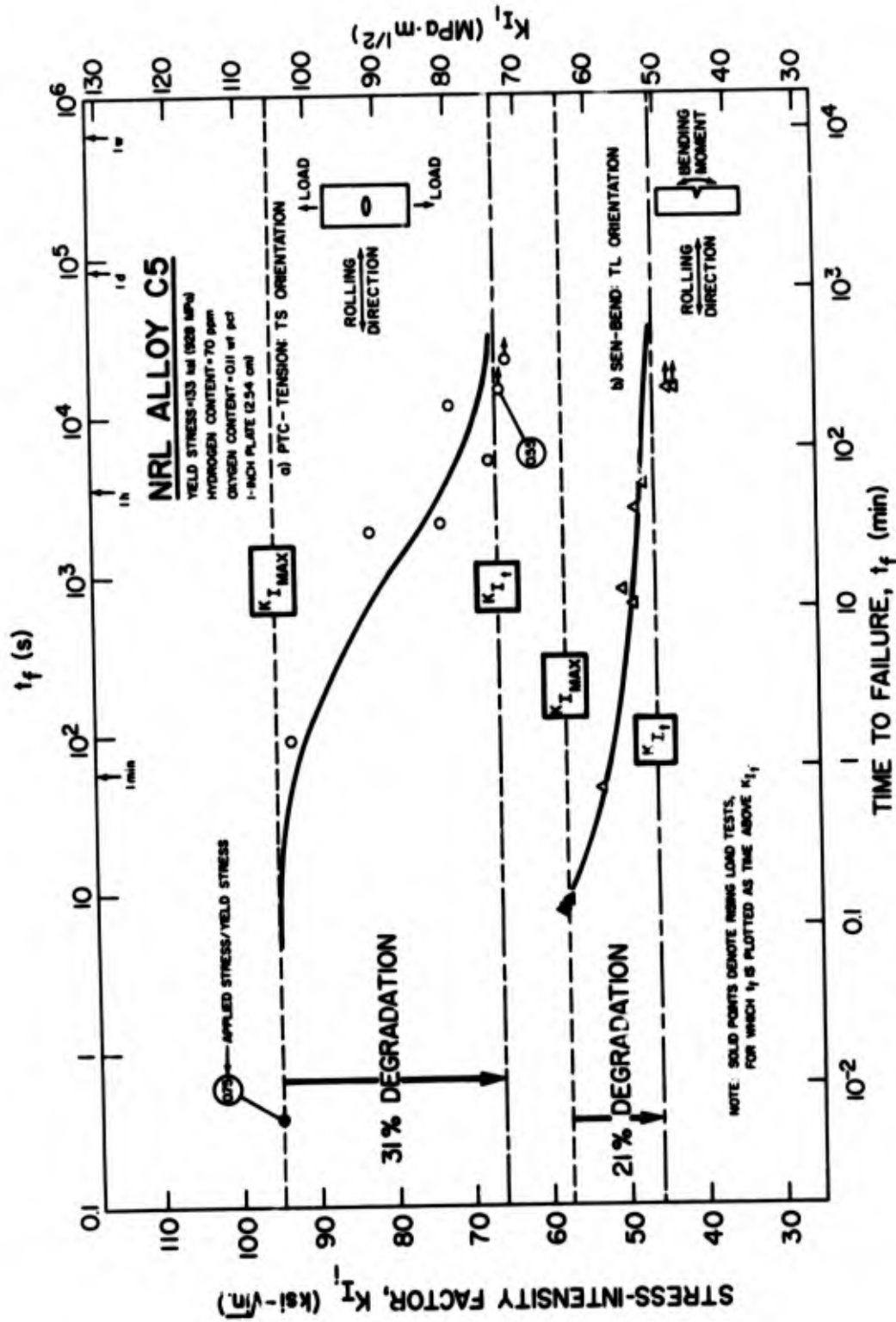


Fig. 3—Sustained-load cracking in NRL alloy C5 in ambient, room-temperature air:
 (a) PTC tension specimens, (b) SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

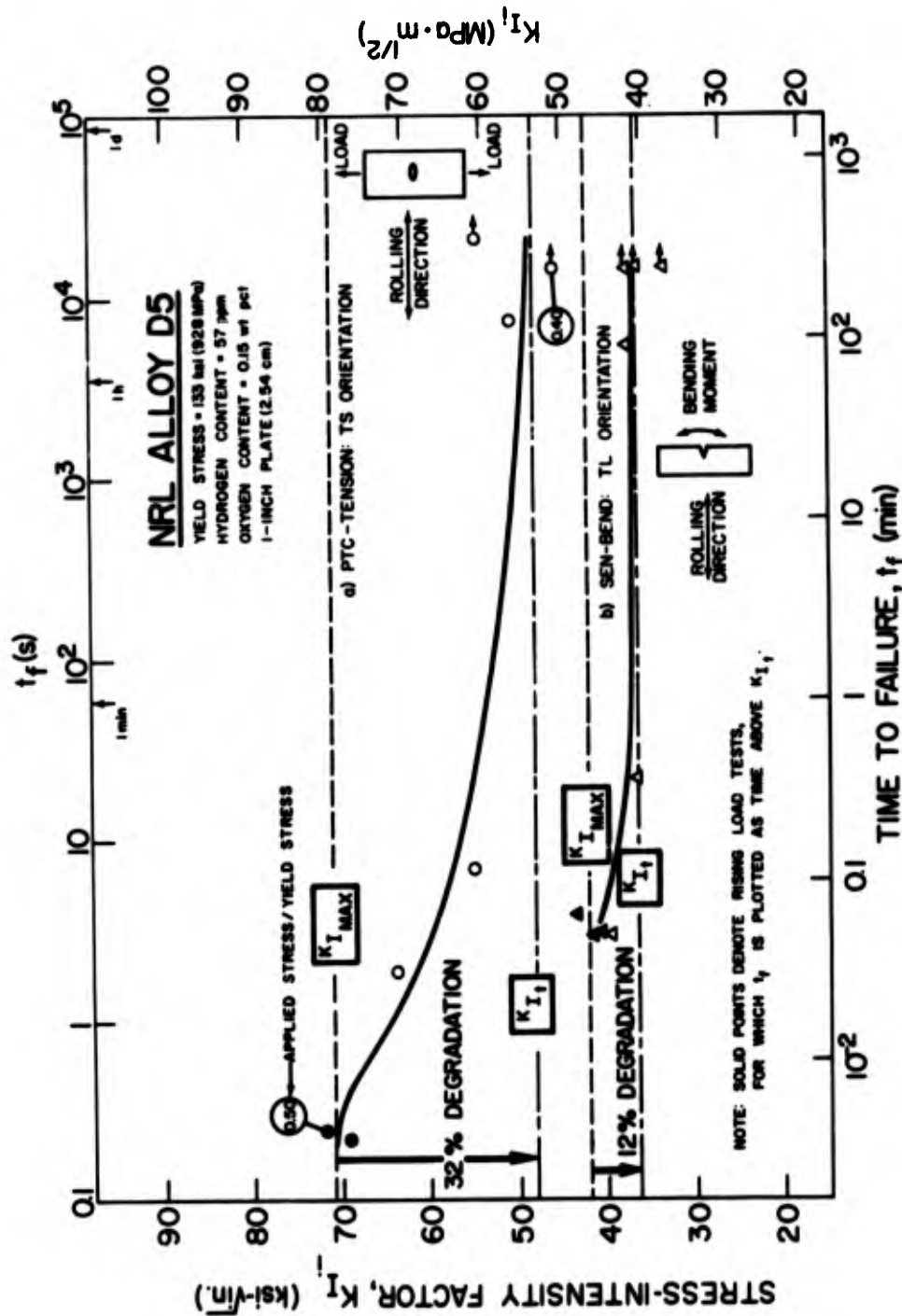


Fig. 4—Sustained-load cracking in NRL alloy D5 in ambient, room-temperature air:
 (a) PTC tension specimens, (b) SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

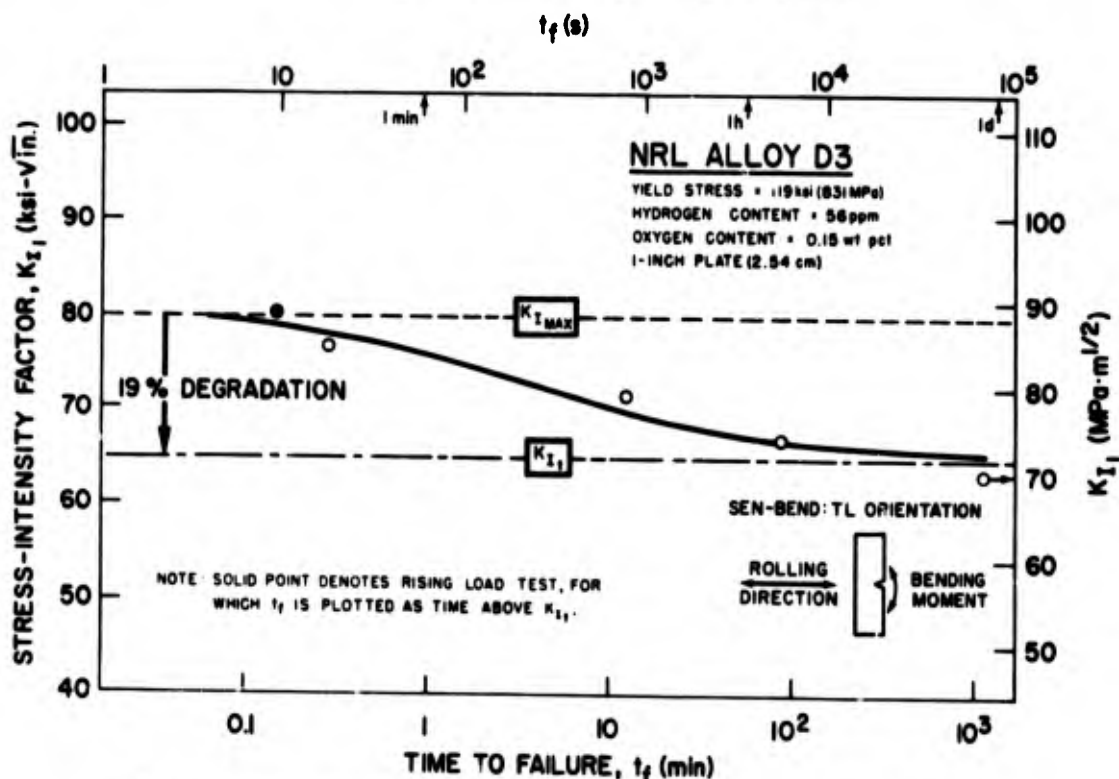


Fig. 5—Sustained-load cracking in NRL alloy D3 in ambient, room-temperature air; SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

corroborate an association between $K_{I\max}$ from the rising-load three-point bend test and the cantilever bend data, a special test was devised using alloy R23A, wherein a rising-load cantilever bend test was performed by employing an Instron machine to drive downward the end of the cantilever bar. As illustrated in Fig. 2, this test indicated that $K_{I\max} = 78 \text{ ksi}\sqrt{\text{in.}}$ ($86.8 \text{ MPa}\cdot\text{m}^{1/2}$), a value which compares favorably with numbers from the rising-load three-point bend tests, viz. $K_{I\max} = 79, 80, 85 \text{ ksi}\sqrt{\text{in.}}$ ($87.9, 89.0, 95.6 \text{ MPa}\cdot\text{m}^{1/2}$). Therefore, $K_{Ir} = K_{I\max}$ is chosen for the computation of K_{It}/K_{Ir} in Eq. (3), as well as for the value of K_{Ii} displayed in Figs. 2-9 for rising-load test results.

Whereas every material tested exhibited degradation in load-carrying capacity owing to SLC, with Δ ranging from about 11 to 35%, for a given crack orientation Δ does not seem to correlate with other variables listed in Table 1, including hydrogen content.

A consideration of Figs. 2-4 (or Table 1) for comparison of results from plates tested in both the SEN and PTC specimen configurations, viz. alloys C5, R23A, and D5, shows that there are large discrepancies in the data for each material between threshold K_{It} values obtained from the two specimen types; similarly for the $K_{I\max}$ values. For example, in the case of alloy C5, $K_{It} = 66$ and $46 \text{ ksi}\sqrt{\text{in.}}$ (73.4 and $51.2 \text{ MPa}\cdot\text{m}^{1/2}$), while $K_{I\max} = 95$ and $58 \text{ ksi}\sqrt{\text{in.}}$ (105.7 and $64.5 \text{ MPa}\cdot\text{m}^{1/2}$) for the PTC and SEN specimens, respectively; the consequent discrepancy in Δ is 31 vs 21%. These differences are probably attributable to anisotropy, since different crack orientations were used for the two specimen types. However, they may also in part be due to the invalid nature of the fracture toughness numbers, although for alloy D5, K_{It} values from both the PTC and SEN specimens satisfy the ASTM thickness requirement. The macrofractograph in

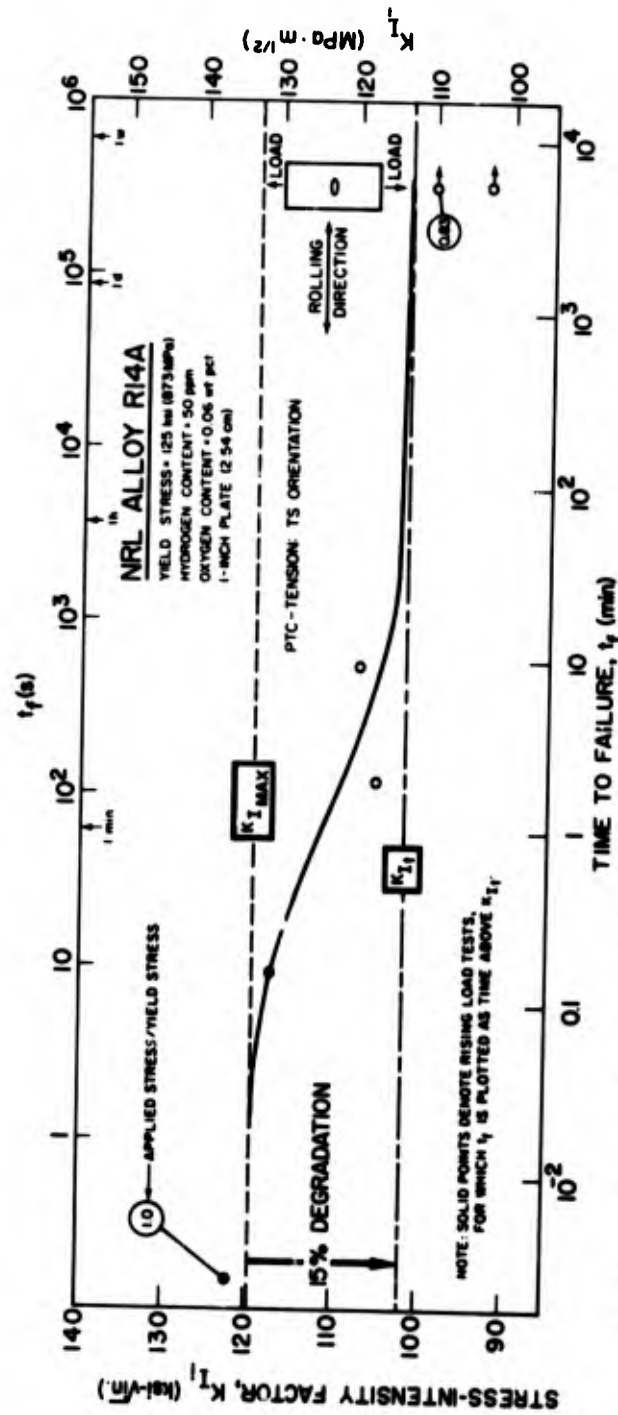


Fig. 6—Sustained-load cracking in NRL alloy R14A in ambient, room-temperature air; PTC tension specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

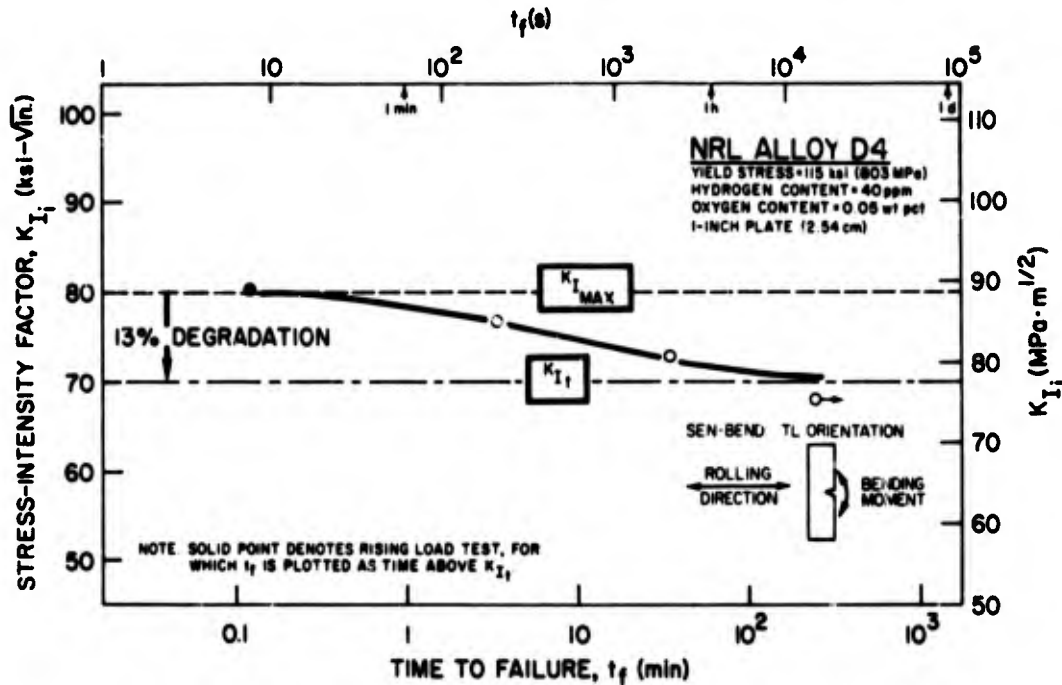


Fig. 7—Sustained-load cracking in NRL alloy D4 in ambient, room-temperature air; SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

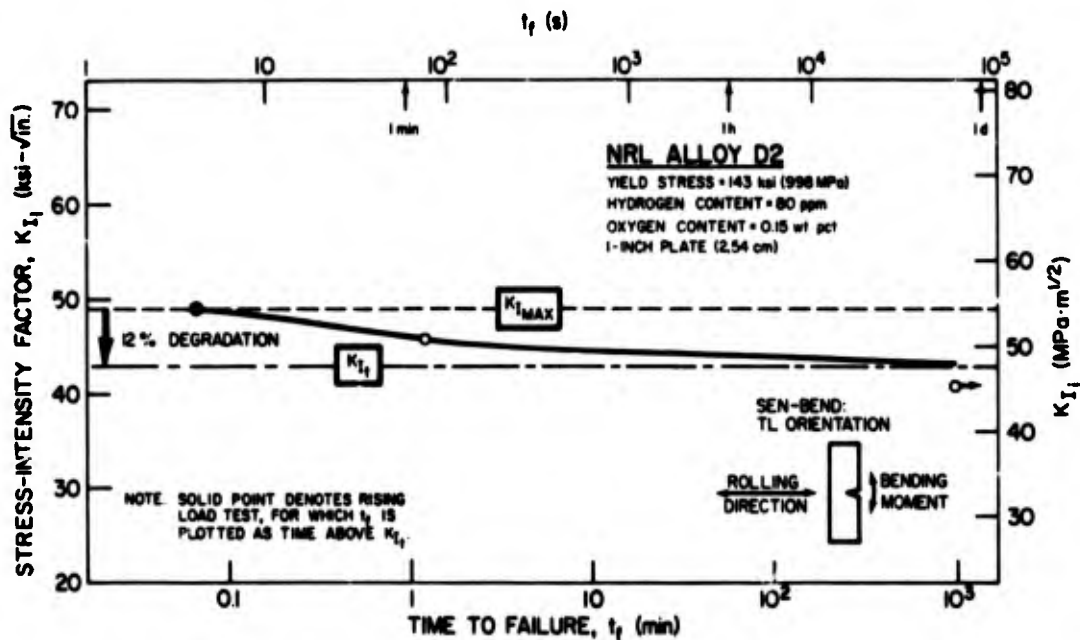


Fig. 8—Sustained-load cracking in NRL alloy D2 in ambient, room-temperature air; SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

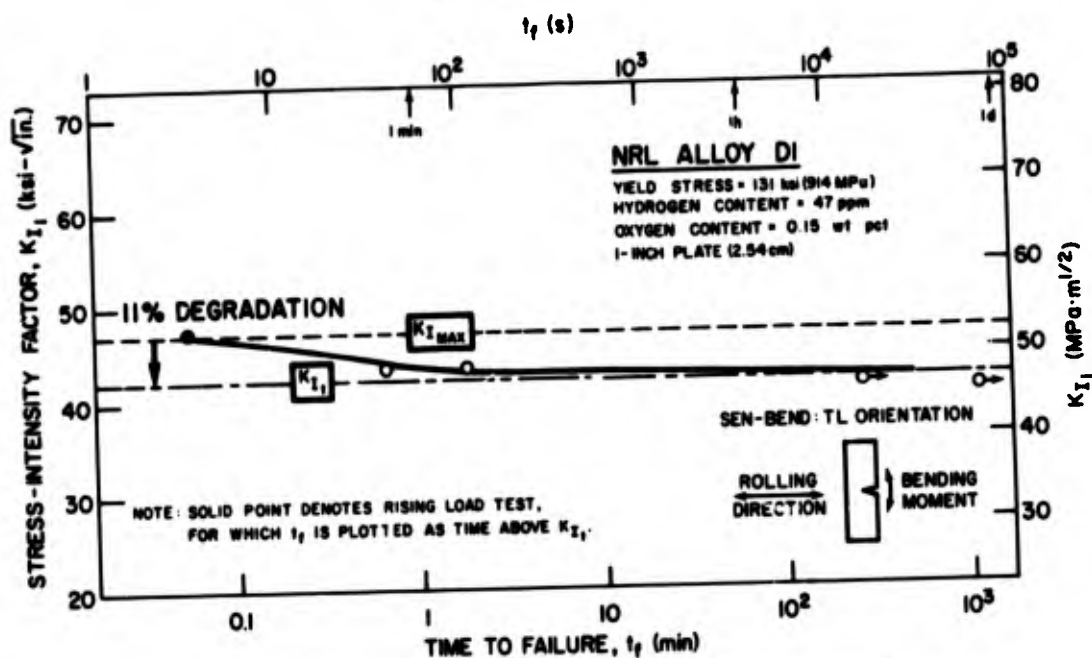


Fig. 9—Sustained-load cracking in NRL alloy D1 in ambient, room-temperature air; SEN bend specimens. Solid points represent rising-load tests; hollow points denote sustained-load tests.

Fig. 10 illustrates the anisotropy in SLC growth in a PTC specimen of alloy R23A. Growth in the region between dashed lines is less in the thickness direction (indicated by arrow) than in the transverse direction, although analysis of the maximum Mode I stress-intensity factor for a given applied load would predict greater growth in the thickness direction for an isotropic material. Scanning electron microscopy shows that SLC growth occurs in alloy R23A by both microvoid coalescence and cleavage. In the SEN bend specimen loaded to an initial $K_{I1} = 65 \text{ ksi}\sqrt{\text{in.}}$ ($72.3 \text{ MPa}\cdot\text{m}^{1/2}$), cleavage was found in decreasing amounts as the crack advanced under sustained load. An example of a cleaved region embedded in a matrix of microvoid coalescence in this specimen is shown in Fig. 11.

Ratios of applied stress to yield stress are encircled in figures containing PTC specimen data; these imply directly the reductions in load-carrying capacity owing to SLC. For example, in the case of alloy R23A, the fracture resistance inferred from the rising-load tests suggests that the fracture-safe, load-carrying limit is approximately 90% of the yield limit load, while fracture resistance implied from the threshold K_{I1} indicates a fracture-safe, load-carrying limit which is only 70% of the yield limit load.

DISCUSSION

While variations in Δ from plate to plate, for a given crack orientation, do not seem to correlate with differences in hydrogen content (among other parameters), it must be cautioned that invalidity of the fracture toughness numbers (cf. Eqs. (4) and (5)) makes quantitative comparisons between materials uncertain. The results of Sandoz (5), for example, suggest that the apparent degradation for a given material and crack orientation depends on specimen thickness, i.e., the degree of constraint relaxation. Thus it would be most appropriate to normalize all of the present results to the same level of constraint

Table 1
Properties of Ti-6Al-4V Alloys

NRL Alloy	Specimen Type		Hydrogen ppm	Oxygen ppm	σ_{ys} ksi (MPa)	Rolling Temp. °F (°K)	Forge Temp. Range	K _{Ir}		K _{Ic} Criteria		K _{It} ksi $\sqrt{\text{in.}}$ (MPa - m ^{1/2})	$2.5(K_{It}/\sigma_{ys})^2 \leq B$	SLC Degradation $\Delta(\%)$
	SEN Bend	PTC Tension						K _Q ksi $\sqrt{\text{in.}}$ (MPa - m ^{1/2})	K _I max ksi $\sqrt{\text{in.}}$ (MPa - m ^{1/2})	$P_{max}/P_Q \leq 1.10$	$2.5(K_Q/\sigma_{ys})^2 \leq B, a, D-a$			
D4	X		40	500	115 (803)	1700 (1200)	β	68 (76)	80 (89)	no	no	70 (78)	no	13
C5	X	X	70	1100	133 (928)	1700 (1200)	$\alpha+\beta$	56 (62)	58 (65) 95 (106)	yes	yes	46 (51) 66 (73)	yes yes	21 31
D3	X		56	1500	119 (831)	1900 (1311)	β	73 (81)	80 (89)	yes	no	65 (72)	yes	19
R23A	X	X	35	1600	144 (1005)			66 (73)	82 (91) 121 (135)	no	no	53 (59) 86 (96)	yes no	35 29
D2	X		80	1500	143 (998)	1400 (1033)	β	48 (53)	49 (55)	yes	yes	43 (48)	yes	12
D1	X		47	1500	131 (914)	1700 (1200)	β	46 (51)	47 (52)	yes	yes	42 (47)	yes	11
D5	X	X	57	1500	133 (928)	1700 (1200)	$\alpha+\beta$	41 (46)	42 (47) 71 (79)	yes	yes	37 (41) 48 (53)	yes yes	12 32
R14A		X	50	600	125 (873)				120 (133)			102 (113)	no	15

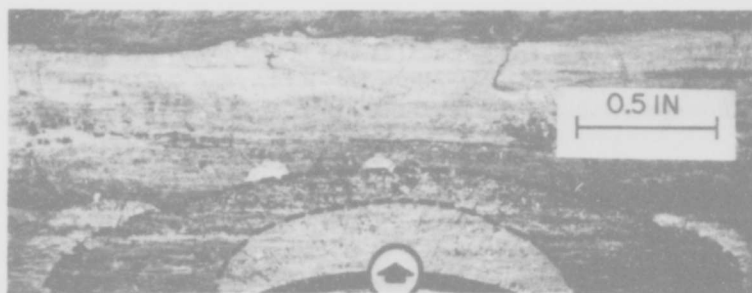


Fig. 10—Anisotropic crack growth in a PTC tension specimen of NRL alloy R23A under sustained load: SLC growth (bounded by dashed line) is less in thickness direction (denoted by arrow) than in transverse direction.

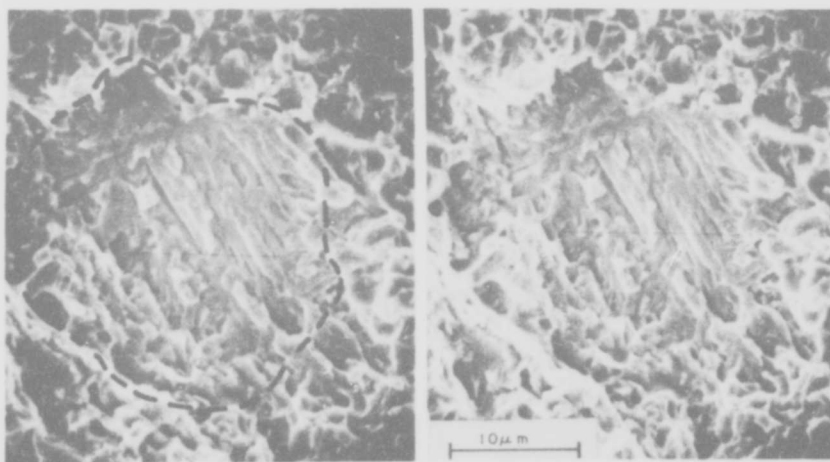


Fig. 11—Stereoscopic pair of scanning electron micrographs illustrate cleaved region (within dashed loop) embedded in matrix of coalesced microvoids, from SEN bend specimen of NRL alloy R23A, cracked under sustained load with $K_{Ii} = 65 \text{ ksi}\sqrt{\text{in.}}$ ($74 \text{ MPa}\cdot\text{m}^{1/2}$)

(or relaxation thereof); unfortunately, fracture mechanics has yet to be developed to this capability.

However, most of the threshold K_{It} values do meet the ASTM thickness requirement

$$2.5 \left(\frac{K_{It}}{\sigma_{ys}} \right)^2 \leq B \quad (6)$$

for validity, and thus may be useful for fracture mechanics interpretations of flaw size vs stress level relationships. Hence the SLC effect may be reckoned with in much the same

manner that stress-corrosion-cracking effects are factored into fracture-safe design procedures, as described by Judy and Goode (1) and Pellini (16). For plates of those materials which do not satisfy Eq. (6), viz. alloys D4, R14A, and R23A (PTC specimen, TS orientation), flaw size vs stress level relationships independent of thickness cannot be inferred.

Perhaps it is worth emphasizing that a fracture toughness number K_{It} in Table 1 which satisfies Eq. (6) appears to represent a more conservative fracture-safe design parameter than a K_{Ic} number, determined from a rising-load test, in accordance with ASTM specifications, which cannot account for insidious time-delayed failures owing to SLC. For example, in the case of alloy C5 (SEN specimen data) the K_Q value of $56 \text{ ksi}\sqrt{\text{in.}}$ ($62.3 \text{ MPa}\cdot\text{m}^{1/2}$), which is a close approximation to K_{Ic}^* , implies a fracture resistance some 21% higher than the threshold $K_{It} = 46 \text{ ksi}\sqrt{\text{in.}}$ ($51.2 \text{ MPa}\cdot\text{m}^{1/2}$) determined from sustained-load, cantilever bend tests.

To investigate any potential role which hydrogen may have in SLC, an analysis of crack growth rate data might prove useful in detecting an incubation time τ (7, 17, 18) associated with this cracking. Such a τ should exist if a critical concentration of hydrogen c must accumulate at the crack tip before cracking can proceed. Hydrogen might so accumulate by diffusing from the bulk by interaction with the crack-tip stress field as modeled by Liu (19), with the result that

$$c = c_0 \exp \left(\frac{AK_{II}}{kT\sqrt{r}} \cos \frac{\theta}{2} \right), \quad (7)$$

where

c_0 = concentration of hydrogen in the bulk

A = lumped material constant

k = Boltzmann's constant

T = absolute temperature

r, θ = polar coordinates emanating from crack tip.

By using the compliance technique (2), crack growth rate data shown in Fig. 12 have been obtained from the SEN specimen of alloy D3, which was loaded initially to $K_{II} = 66.6 \text{ ksi}\sqrt{\text{in.}}$ ($74.1 \text{ MPa}\cdot\text{m}^{1/2}$) and failed under sustained load in room air after about 1.5 hr. Clearly the data in Fig. 12 do not reveal an incubation period, as crack growth begins almost immediately, at rates of the order of 10^{-5} in./s (10^{-5} cm/s). If the high mobility of interstitial hydrogen has made the incubation period undetectably small at room temperature, then lower temperatures can be used to increase the extent of any incubation period, which should vary inversely with drift velocity of the hydrogen. The drift velocity used by Cottrell and Bilby (20, 21) in relating solute interaction with a dislocation stress field is given by Einstein's theory of Brownian motion as

*Supplementary tests with non-sidegrooved specimens, with $D = 2.0 \text{ in.}$ (5.08 cm), indicated that $K_{Ic} = 55.4 \text{ ksi}\sqrt{\text{in.}}$ ($61.6 \text{ MPa}\cdot\text{m}^{1/2}$), as determined in strict accord with ASTM procedures. Similarly, for alloy D3, $K_{Ic} = 71.8 \text{ ksi}\sqrt{\text{in.}}$ ($79.9 \text{ MPa}\cdot\text{m}^{1/2}$), which compares well with the value K_Q reported in Table 1, $73 \text{ ksi}\sqrt{\text{in.}}$ ($81.2 \text{ MPa}\cdot\text{m}^{1/2}$).

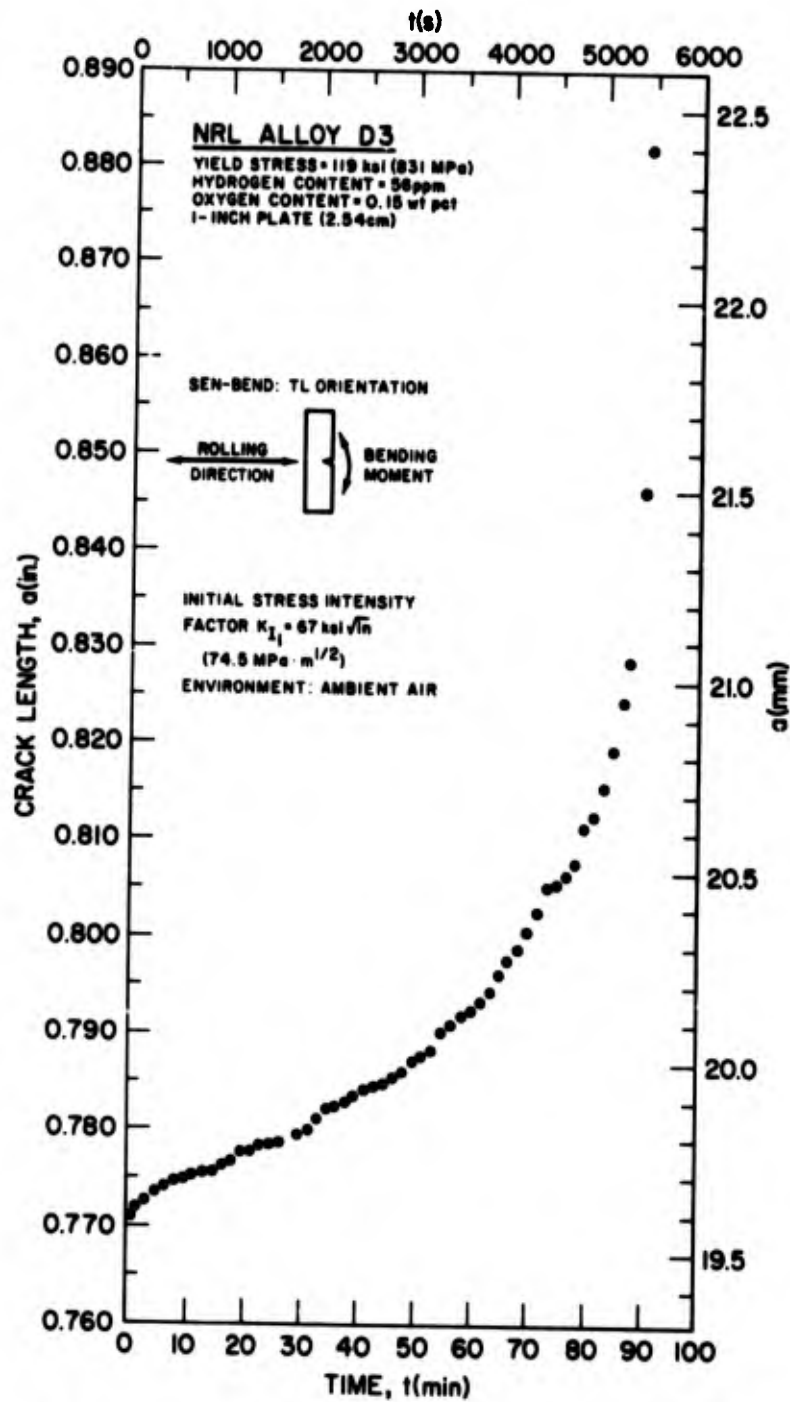


Fig. 12—Crack extension in a SEN bend specimen of NRL alloy D3 under sustained load with $K_{I1} = 67 \text{ ksi}\sqrt{\text{in}}$ (74 MPa·m^{1/2})

$$\bar{v} = - \left(\frac{D}{kT} \right) \nabla V, \quad (8)$$

where D is the diffusivity and V the interaction energy. Liu (19) expresses the interaction energy between the diffusing hydrogen and the stress field of the crack tip as

$$V = - \frac{AK_{II}}{\sqrt{r}} \cos \frac{\theta}{2} . \quad (9)$$

From Eqs. (8) and (9) it follows that the incubation time is given by

$$\tau = \frac{2LkT\tau^{3/2}}{AK_{II} D_0 \exp \left(\frac{-Q}{kT} \right)} , \quad (10)$$

where

L = a constant with the dimension of length

D_0 = the diffusion constant

Q = the activation energy.

At this point, however, any participation of hydrogen in the SLC process remains undefined.

Furthermore, a recent study by Meyn (22) showed that for one Ti-6Al-4V alloy the SLC effect failed to disappear at an exceedingly low hydrogen level (i.e., a few ppm) which bordered on the limit of detection. Thus it is not clear that SLC in titanium alloys is caused solely by internal hydrogen content. In other studies of SLC in inert environments, it has been noted that crack growth occurs with an initial region of transient growth, followed by a region of steady state growth, and terminating with a region of accelerating growth which leads to failure; this appears to be the pattern in Fig. 12. Such behavior has in some instances been relatable to material flow properties (23). Similar efforts might prove fruitful in the present case to interpret room-temperature crack growth rate data.

SUMMARY

Degradation in load-carrying capability owing to SLC is widespread and serious in alloys of the Ti-6Al-4V family. Each of eight plate samples tested in air at room temperature exhibited the effect, with degradations ranging from 11 to 35%. The amount of degradation does not seem to correlate with interstitial contents, processing variables, or strength and toughness levels; however, the susceptibility to SLC is orientation dependent. For most of the materials examined, invalidity of the rising-load fracture toughness values with respect to plane-strain criteria makes quantitative comparison of degradations uncertain.

If the potential hazard of time-delayed catastrophic failures owing to SLC is to be avoided in structures fabricated with these alloys, the SLC effect is one that must be reckoned with, in much the same manner that stress-corrosion-cracking effects are factored into fracture-safe design procedures. It appears for most materials examined that the threshold stress-intensity factor K_{II} , below which time-delayed failures owing to SLC do not occur, may be useful for fracture mechanics interpretations of flaw size vs stress level relationships. While there is no standard technique for determining such a threshold,

it appears that K_{It} represents a more conservative, fracture-safe design parameter than K_{Ic} , determined from a rising load test in conformity with ASTM specifications, which cannot account for insidious time-delayed failures owing to SLC.

The mechanism of SLC has yet to be defined, especially with regard to the possible role of interstitial hydrogen.

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