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Mechanical Behavior, Microstructure and Thermooxidation Properties of Sequentially Crosslinked Ultrahigh Molecular Weight Polyethylenes

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ABSTRACT: The aim of this study was to explore the impact of the sequential irradiation and annealing process on the microstructure, thermooxidation behavior and mechanical properties of GUR 1050 ultrahigh molecular weight polyethylene (UHMWPE) with respect to the postirradiation annealed material. For this purpose, the effects of a variety of irradiation and annealing conditions on microstructure and mechanical properties were investigated. Differential scanning calorimetry was performed to characterize melting temperature, crystalline content and crystal thickness, whereas transmission electron microscopy provided additional insights into crystal morphology. Thermogravimetric experiments in air served to assess thermooxidation resistance and changes associated to radiation-induced crosslinking. Fatigue properties were studied from three different approaches, namely short-term cyclic stress–strain tests, long-term fatigue experiments and crack propagation behavior. Likewise, three experimental techniques (uniaxial tensile test, impact experiments, and load to fracture of compact tension specimens) allowed evaluation of the fracture resistance. The present findings confirm sequentially crosslinked UHMWPE exhibited improved thermooxidation resistance and thermal stability compared to post-irradiation annealed UHMWPE. Also, the mechanical behavior, including the fatigue and fracture resistance, of these materials was generally comparable regardless of the annealing strategy. Therefore, the sequential irradiation and annealing process might provide higher oxidation resistance, but not a significant improvement in mechanical properties compared to the single radiation dose and subsequent annealing procedure. © 2013 Wiley Periodicals, Inc. J. Appl. Polym. Sci. 129: 2518–2526, 2013

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INTRODUCTION

Nowadays, modern highly crosslinked polyethylenes (HXLPE) have replaced conventional, gamma inert sterilized, ultrahigh molecular weight polyethylene (UHMWPE) for use in total hip arthroplasties.[1,2] The rationale behind the introduction of HXLPE for orthopedic use is a dramatically improved wear resistance, which, in turn, stems from the elevated crosslink density that high gamma or electron beam radiation doses (~ 100 kGy) impart UHMWPE.[3–5] This beneficial property has made possible a significant reduction in the incidence of revisions that historical, gamma air sterilized, and conventional UHMWPE inserts experienced due to osteolytic reactions and eventual aseptic loosening triggered by UHMWPE wear debris particulate.[6] Despite its positive effect on wear resistance, irradiation unavoidably generates free radicals, which have the potential to initiate the oxidation cycle of HXLPE. To prevent long-term degradation in the presence of oxygen, postirradiation thermal treatments have been necessary to eliminate, or at least reduce, radiation-induced free radicals.[7] In general, thermal treatments used in first generation HXLPE production can be classified as annealing or remelting depending on whether or not they were conducted below the melting temperature. Despite the excellent wear resistance of current HXLPE, they present some drawbacks. On one hand, annealed HXLPE contain residual free radicals, and therefore *in vivo* oxidation may result in material embrittlement, ultimately compromising the mechanical performance of the insert. High radiation doses followed by remelting, on the other hand, considerably reduce the fatigue and fracture properties of UHMWPE.[8–12] Although the clinical performance of HXLPE in total hip arthroplasty has been satisfactory for the first decade of use,[6] there is growing evidence of high oxidation in nonload-bearing regions of

retrieved annealed hip inserts,[13] and early crack initiation in few, case studies, remelted retrievals.[14,15] With regard to total knee arthroplasty, annealed and remelted HXLPE are not generally recommended, since *in vivo* oxidation, and rapid cracking, respectively, might have dramatic consequences under the more demanding conditions of the knee joint.[16]

Second-generation HXLPE represent an attempt to simultaneously provide oxidative stability and preserved mechanical properties. Three main strategies have been proposed, namely sequential irradiation and annealing, incorporation of antioxidants (i.e., vitamin E) by blending or diffusion, and mechanical annealing. Vitamin E acts as a scavenger of radiation-induced free radicals in UHMWPE,[17–19] whereas solid-state deformation below the melting point of HXLPE provides enhanced strength and good oxidative stability.[20] The basis for sequentially irradiated and annealed UHMWPE is that the annealing treatment would be more effective eliminating free radicals produced by 30 kGy-irradiation steps, since chain mobility is higher when crosslink density is low. Thus, three consecutive irradiation and annealing cycles would provide the excellent wear resistance associated to high radiation doses ($\sim$ 100 kGy) as well as oxidative stability, without negatively affecting crystallinity and mechanical properties.[21–23] Terminal gas plasma sterilization completes the production of commercial sequentially irradiated and annealed UHMWPE. However, some contradictory results appear in the literature concerning the improvement in mechanical properties obtained with this new sequential irradiation and annealing process.[23–25]

In the context presented above, the current work aims at comparing several second generation, sequentially irradiated and annealed UHMWPEs, with first generation, postirradiation annealed, HXLPE from mechanical, and thermooxidation perspectives. The correlation between microstructural features and mechanical properties was also studied.

MATERIALS AND METHODS

Materials

We used GUR 1050 UHMWPE in the form of compression-molded sheets (Orthoplastic Medical; Lancashire, UK) as raw material. Crosslinking was achieved by three consecutive 30 kGy gamma irradiation in air steps (Aragogamma S.A.; Barcelona, Spain). Postirradiation annealed UHMWPE was obtained performing a terminal annealing treatment at 130°C for 8 h in a vacuum oven (Weiss-Gallenkamp; Loughborough, UK). Sequentially irradiated and annealed UHMWPEs were produced conducting identical annealing treatments after all or some of the 30 kGy irradiation steps. A final machining step was necessary to remove the outer, oxidized, layer ($\sim$ 2–3 mm) of the irradiated and annealed preforms, thus obtaining mechanical specimens ready to test. Typically, we used unirradiated UHMWPE as control material, but in some cases also single-dose (90 kGy) irradiated and three-step irradiated (30–30–30) UHMWPEs without further annealing, to discriminate the separate effects of irradiation and annealing. Thus, the seven material groups studied will be referred to as virgin or unirradiated, G0 (30–30–30), G1 (30–30–30A), G2 (30–30A–30A), G3 (30A–30A–30A), G4 (30A–30–30), and G5 (90), where G, 30, and A stand for gamma irradiated, a single 30 kGy irradiation step, and annealing, respectively. It is worth noting that G1, and G3 were obtained following procedures similar to those used to produce commercially available postirradiation annealed HXLPE, and the second generation, sequentially crosslinked HXLPE, respectively.

Differential Scanning Calorimetry and Thermogravimetry

Differential scanning calorimetry (DSC) experiments were conducted in air using a Dynamic Scanning Calorimeter (TA Instruments Q20). At least three samples ($n \geq 3$) per material group were heated from room temperature to 200°C at a 10°C/min rate. The area below the first-heating DSC curves from 80 to 160°C, normalized by 290 J/g as the enthalpy of melting of a 100% crystalline polyethylene, served to calculate crystallinity contents. The melting transition temperature was registered as the peak temperature of the melting endotherm.

Thermogravimetric (TG) experiments were carried out in air using a TA Instrument Q5000 thermobalance (accuracy: 10^{-4} mg). Six milligram-samples ($n \geq 3$ per material group) were heated from room temperature to 800°C at a 10°C/min rate. The main features in decomposition curves were documented and analyzed following recently reported guidelines.[26]

Transmission Electron Microscopy analysis

Transmission electron microscopy (TEM) was performed to observe the microstructure of all the materials using a Jeol 100CX microscope operating at 100 kV. Appropriate specimen preparation involving clorosulphonic staining of thin films was necessary to make the samples ready for TEM, as reported elsewhere.[27] 20,000×, 40,000× and 60,000× magnification micrographs were taken and analyzed using Digital Micrograph 3.3.1 (Gatan, Pleasanton, CA) to measure changes in lamellar thickness after irradiation and annealing stabilization.

Uniaxial Tensile Tests

Uniaxial tensile tests ($n \geq 3$) per ASTM D638 were performed in an electromechanic Instron machine on type M-I specimens, at $T = 23 \pm 2$°C, with a displacement rate of 5 mm/min (initial nominal strain rate $\sim 0.002\ s^{-1}$). In addition to typical parameters, such as yield stress, elastic modulus, and ultimate stress and strain, work to fracture values were calculated from engineering stress–strain plots.

Cyclic Stress–Strain, Long-Term Fatigue, and Fatigue Crack Propagation Experiments

The fatigue behavior was characterized by means of three experimental techniques. First, cyclic stress–strain experiments were conducted on tensile specimens for up to 50 cycles. A displacement rate of 15 mm/min (initial nominal strain rate 0.005 s^{-1}), and a maximum nominal stress, σ_{max}, of 16 MPa (stress ratio $R \sim 0$) were chosen to conduct these experiments. At least three samples ($n \geq 3$) per material group were tested in an Instron 5565 machine at 24 ± 1 °C. These short-term cyclic experiments provided information about the total plastic strain reached, $\varepsilon(50)$, and the secant modulus at the first cycle, $E_s(1c)$, which are measures of the material softening, and stiffness, respectively.

Second, long-term fatigue tests, *S*/*N* stress-life experiments, were performed on dog bone specimens using a servohydraulic Instron 8032 machine and following ASTM E606 guidelines. These tests ran under load control following a sine waveform (frequency 1 Hz; stress ratio $R \sim 0$). The strain was continuously monitored employing an extensometer and the testing temperature was 23 ±

Table I. Thermal and Thermogravimetric Parameters (Mean ± Standard Deviation) Obtained from DSC and TG Experiments for Unirradiated, Postirradiation Annealed, and Sequentially Crosslinked UHMWPEs

	DSC			TG		
Material	Shoulder temperature (°C)	Melting temperature (°C)	Crystallinity (%)	T_B (°C)	T_0 (°C)	T_1 (°C)
Unirradiated	N/A	136.2 ± 0.3[a]	51.9 ± 0.7[a]	151.9 ± 0.6[a]	214.9 ± 0.6[a]	374.4 ± 3.8[a]
G0 (30-30-30)	N/A	142.1 ± 1.1[b]	58.7 ± 0.5[b]	140.7 ± 0.1[b]	219.3 ± 0.4	388.0 ± 2.4[b,c]
G1 (30-30-30A)	125.5 ± 1.6	141.0 ± 0.6[b]	53.1 ± 1.0[c]	142.8 ± 3.4[b]	212.6 ± 0.9[b]	384.3 ± 4.7[b]
G2 (30-30A-30A)	126.0 ± 0.2	141.1 ± 0.6[b]	51.6 ± 1.1[c]	144.4 ± 1.1[b]	211.8 ± 1.7[b]	390.7 ± 3.6[b]
G3 (30A-30A-30A)	127.5 ± 0.3	141.4 ± 0.5[b]	53.7 ± 2.1[c]	167.6 ± 2.5[c]	233.0 ± 0.8[c]	397.6 ± 5.0[b,d]
G4 (30A-30-30)	N/A	143.2 ± 0.9[b]	58.8 ± 0.6[b]	141.2 ± 0.8[b]	216.2 ± 1.9	391.5 ± 1.2[b]
G5 (90)	N/A	141.1 ± 0.5[b]	58.8 ± 0.2[b]	139.9 ± 0.4[b]	217.8 ± 3.5	396.1 ± 12.2[b]
		$P < 0.0001$[a,b]	$P < 0.0001$[a,b]	$P \leq 0.003$[a,b]	$P \leq 0.03$[a,b]	$P \leq 0.023$[a,b]
			$P < 0.0001$[b,c]	$P < 0.0001$[a,c; b,c]	$P < 0.0001$[a,c; b,c]	$P < 0.0054$[c,d]

N/A: Not applicable.
[a, b; a, c; b, c; c, d]Levels of significance (P-values) are given for statistical comparisons between material properties labeled with the corresponding superscripts.

2°C. The selected failure criterion was the number of cycles needed to reach a 12% strain level, as in previous studies.[9] This strain level is close to strain maxima (12–15%) registered in UHMWPE tibial components, and has been connected to the appearance of fatigue-induced microscopic defects.[9,28,29]

Third, near-threshold fatigue crack propagation (FCP) experiments were performed on standard compact tension specimens per ASTM E647. All compact specimens were precracked using a razor blade. A digital camera allowed crack growth monitoring and gradual crack length assessments. At least three specimens ($n \geq 3$) were tested per material group and they underwent tension cycling (frequency 5 Hz) with $R = 0.1$. $\Delta K_{inception}$ values and Paris coefficients were obtained from crack propagation curves.

Toughness Characterization

Impact Izod tests ($n \geq 3$ per material group) were carried out at 23 ± 2°C on double-notched specimens following ASTM F648 guidelines. Conversely, work to fracture values obtained as the area below the engineering stress–strain curves of tensile experiments gave an estimation of toughness in a quasi-static situation. Finally, compact tension specimens with dimensions complying ASTM D6068-02 (width 20 mm, thickness 10 mm, and original crack length 10 mm) were loaded to fracture in an attempt to additionally characterize the toughness behavior. Load–displacement curves were registered and analyzed to obtain relevant data.

Statistical Analysis

Student's *t*-tests served to detect significant differences between the thermal, TG, and mechanical properties of the studied material groups. A level of $P < 0.05$ was selected as indicative of significance.

RESULTS AND DISCUSSION

DSC and Thermogravimetry

Both postirradiation annealing and sequential irradiation-annealing caused important changes in the microstructure of UHMWPE. Thus, DSC curves revealed irradiation was responsible for significant 5°C and 12% raises in melting temperature and crystallinity, respectively, as compared with virgin UHMWPE (Table I). Both postirradiation and sequential annealing treatments did not affect melting temperature of as-irradiated UHMWPE, but took crystalline contents back to the level (52–54%) of the unirradiated polymer. Also, materials subjected to two or more irradiation-annealing cycles, G2 and G3, developed an additional endothermic peak at about the annealing temperature (~ 126°C; Figure 1). In contrast, postirradiation annealed UHMWPE, G1, only exhibited a small shoulder at the same temperature. Finally, no significant differences ($P > 0.05$) were detected between the melting temperature and crystallinity of both as-irradiated UHMWPE materials, G0 and G5. The present findings appear to be consistent with the

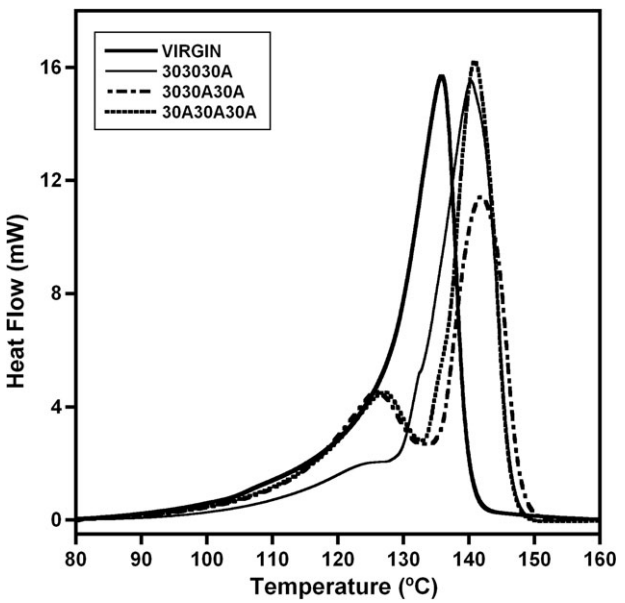

Figure 1. First-heating DSC curves corresponding to virgin, postirradiation annealed (303030A or G1), and sequentially crosslinked (3030A30A or G2, and 30A30A30A or G3) UHMWPEs.

occurrence of radiation-induced recrystallization as proposed by Premnath et al.,[30] on one hand, and partial melting, lamellar thickening and crystallization of smaller crystals during annealing.[31,32] Together, these phenomena would explain the elevated melting temperature, and the appearance of an additional, smaller, endothermic peak in the case of sequentially crosslinked materials. First, molecular rearrangements triggered by irradiation allowed secondary recrystallization onto the surface of original lamellae resulting in elevated melting temperature and crystallinity. This radiation-induced crystallinity increase was lost upon 8 h annealing at 130°C, suggesting that partial melting of lamellar crystals prevailed over lamellar thicknening and crystallization of small crystals. The sequential annealing strategy, however, did not imply an accumulative decrease in crystallinity, probably due to comparatively higher chain mobility (i.e., lower crosslink density), which would favor lamellar thickening during the annealing steps.

The thermooxidation behavior of UHMWPE was also clearly affected by crosslinking and annealing processes. All TG decomposition curves showed a very small but detectable mass increase associated with thermooxidation of the polymer followed by an abrupt weight loss due to thermal degradation [Figure 2(A,B)]. The onset of the thermooxidation process, denoted T_B, reflects the susceptibility to oxidation, as reported previously.[26] In this study, both as-irradiated polyethylenes, G0 and G5, exhibited the lowest T_B values ($P < 0.05$), and a significant decrease in T_B was also observed for postirradiation annealed UHMWPE, G1, and for crosslinked materials subjected to one or two sequential irradiation-annealing steps, G2 and G4 ($T_B \sim 141$–144°C; $P \leq 0.003$; Table I). Overall, these data suggest that the former annealing treatments did not succeed in quenching radiation-induced free radicals and, therefore, in providing complete oxidative stability. This was probably because they were not able to eliminate free radicals trapped in crystalline regions, yielding materials with high susceptibility to oxidation. Three irradiation-annealing steps, however, did not result in a T_B decrease, but in a significant shift towards higher temperatures as registered for G3 specimens [$T_B \sim 167$°C; $P < 0.0001$; Figure 2(B)]. Although thermooxidation could not be completely avoided in sequentially crosslinked materials, the significantly delayed weight gain might be indicative of comparatively higher oxidation resistance. In regard to results corresponding to temperatures at maximum weight, T_0, they followed a trend similar to that of T_B data (Table I).

The beginning of the thermal degradation, indicated by T_1 [Figure 2(A)], was also significantly affected by crosslinking and stabilization processes. They provoked a gradual increase from about 375°C to almost 400°C for unirradiated UHMWPE and sequentially crosslinked UHMWPEs, G3, respectively ($P < 0.0002$). Irradiation processes without further annealing steps also resulted in increased thermal stability ($T_1 \sim 390$°C), which was slightly higher in the case of the single-step irradiated UHMWPE (G5). The present TG results are coherent with crosslink density data trends reported in the literature.[5,9] Previous studies have reported that sequentially annealed UHMWPE exhibits higher crosslink density than postirradiation annealed UHMWPE.[23] Crosslinks between polymeric chains are, in turn,

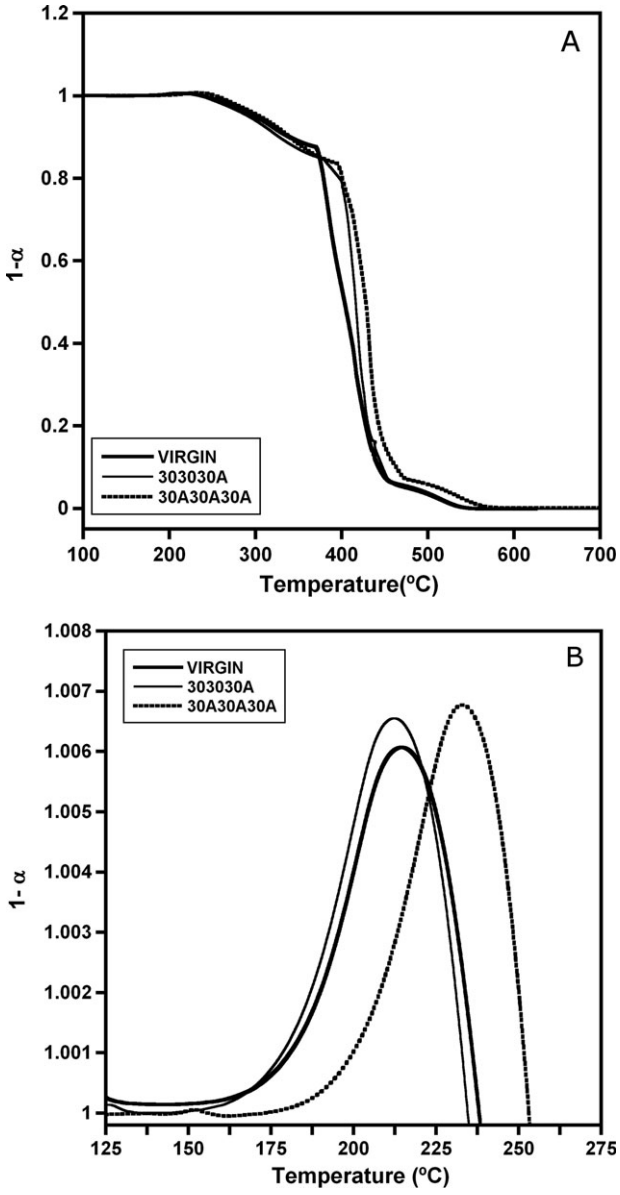

Figure 2. (A and B) TG decomposition curves corresponding to virgin, postirradiation annealed, and sequentially crosslinked UHMWPEs, (A). A close-up view within the 125–275°C range revealed a mass increase associated to thermooxidation of the polymers, (B).

responsible for a concomitant molecular weight increase, and thermal degradation typically begins at increasingly higher temperatures as the molecular weight of the polymer grows.[33] In this sense, researchers have confirmed higher thermal stability, that is higher T_1, of irradiated polyethylenes.[34] Although admittedly the lack of crosslink density assessments in this study impedes to draw definite conclusions, there appears to be a connection between elevated crosslinked density and enhanced thermal stability in orthopedic UHMWPEs.

TEM Analysis

TEM micrographs of unirradiated UHMWPE showed the typical features of a semicrystalline polymer with randomly oriented crystal lamellae immersed in the amorphous region, which

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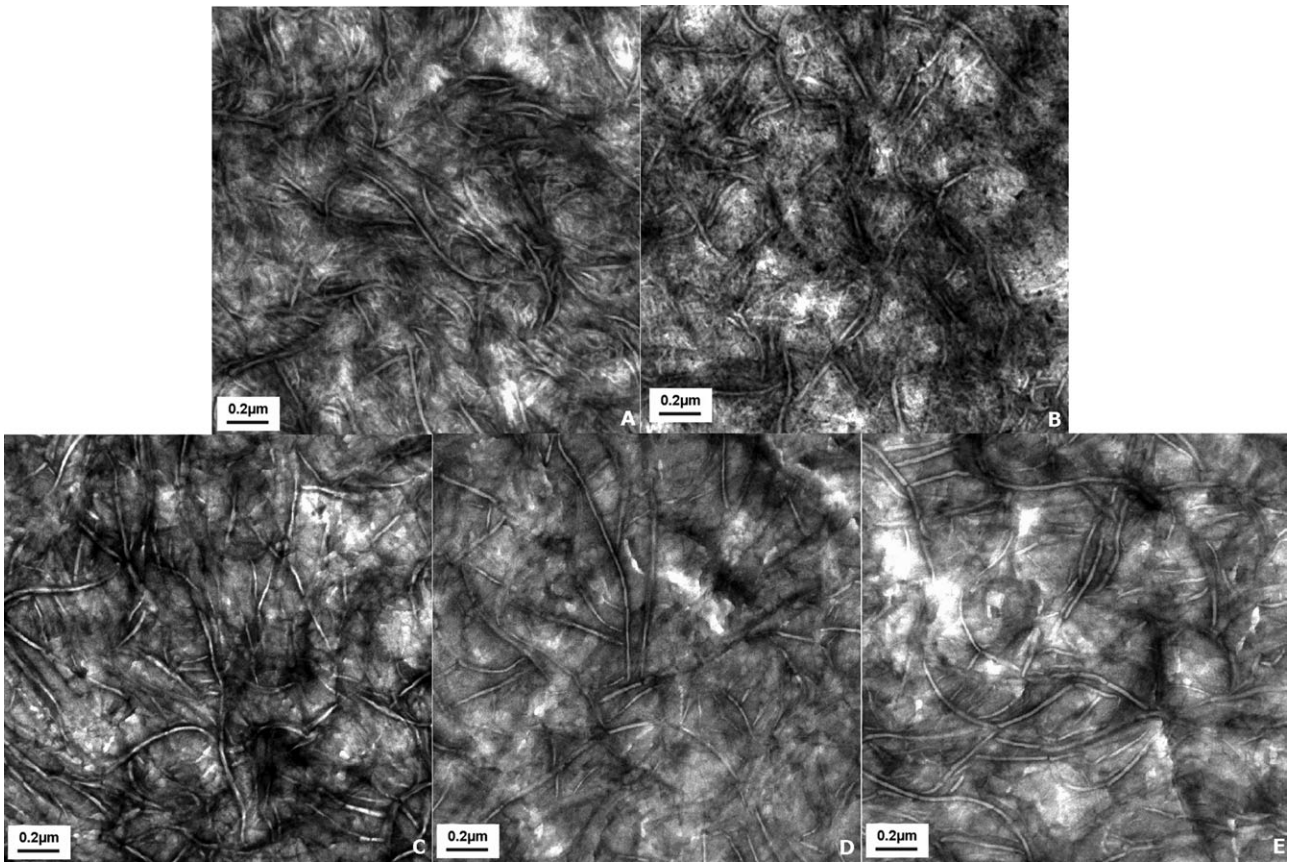

Figure 3. (A–E) TEM micrographs (×40,000) of virgin, (A), as-irradiated (B), postirradiation annealed, (C), sequentially crosslinked G2, (D), and sequentially crosslinked G3, (E), UHMWPEs.

appeared as a dark grey region [Figure 3(A–E)]. The average lamellar thickness of virgin UHMWPE was 29 ± 3 nm, and upon irradiation crystal thickness experienced a small, but statistically significant, increase up to 32 ± 3 nm ($P < 0.0001$). In contrast, the combination of crosslinking and stabilization processes generally resulted in a significant decrease of this property. Sequentially crosslinked, G3, materials were the only exception as they presented crystal thicknesses similar (30 ± 3 nm; $P > 0.42$) to that of unirradiated specimens. The more restricted chain mobility in postirradiation annealed, G1, and G2 UHMWPEs could inhibit crystal thickening during annealing, yielding thinner lamellae (27 ± 2, and 26 ± 3 nm, respectively; $P < 0.0001$ with respect to unirradiated, G0 and G3 UHMWPEs). Lamellar thickening mechanisms, in contrast, would be favored during annealing in sequentially crosslinked materials, G3, due to higher chain mobility.

Uniaxial Tensile Results

Irradiation and stabilization treatments caused considerable changes in mechanical parameters. Both postirradiation annealed and sequentially crosslinked UHMWPEs experienced a strong decrease (∼ 50%) in ductility as reflected by strain to fracture results ($P < 0.0001$; Table II and Figure 4). The fracture stress also decreased after crosslinking and stabilization ($P \leq 0.02$ with respect to unirradiated UHMWPE), but this parameter was almost identical for the three sequentially crosslinked UHMWPEs, G1, G2, and G3 ($P > 0.54$). As-irradiated materials, G0 and G5, exhibited slightly higher yield stresses, fracture stresses, and ultimate strains than sequentially crosslinked materials.

Cyclic Stress–Strain, Long-Term Fatigue, and Crack Propagation Behavior

Cyclic stress–strain experiments confirmed a significant decrease in material softening, $\varepsilon(50)$, upon irradiation (8.3 ± 0.9 and 4.3 ± 0.2 for virgin and as-irradiated, G0, UHMWPEs, respectively; $P < 0.0001$). However, this positive decrease was lost when irradiation and annealing processes were combined. Thus, material softening went down to 7.6 ± 0.8, 9.0 ± 1.4, and 8.5 ± 1.6 for postirradiation, G1, and sequentially annealed, G2 and G3, materials, respectively ($P \leq 0.0008$ with respect to G0 UHMWPE). Likewise, irradiation turned UHMWPE into a stiffer material based on secant modulus results, but, again, the combination of irradiation and annealing processes reverted this change even below the levels of uncrosslinked UHMWPE (Table II).

The combined effects of irradiation and annealing processes caused substantial deterioration of the fatigue strength of unirradiated UHMWPE, regardless of the annealing strategy as shown in the present stress-life, *S–N*, ($S = A \log(N) + B$; A, and B fitting parameters) experiments (Figure 5). In particular, the introduction of annealing treatments between the second and third irradiation steps (i.e., G2 material) did not imply an improvement in long-term fatigue properties, but further

Table II. Mechanical Parameters (Mean ± SD) Obtained from Uniaxial Tension and Cyclic Stress–Strain Experiments for Unirradiated, Postirradiation Annealed, and Sequentially Crosslinked UHMWPE Materials

	Uniaxial Tension			50 Cycles Stress–Strain Testing	
Material	Yield stress (MPa)	Fracture stress (MPa)	Fracture strain	E_S(1c) (MPa)	ε_{16MPa} (50c)
Unirradiated	19.0 ± 0.2	36.3 ± 1.8[a]	8.7 ± 0.5[a]	380 ± 28[a]	8.3 ± 0.9[a]
G0 (30-30-30)	20.8 ± 0.1[a]	36.7 ± 0.3	4.7 ± 0.1[b]	494 ± 13[b]	4.3 ± 0.2[b]
G1 (30-30-30A)	17.7 ± 0.3[b]	30.4 ± 1.8[b]	4.3 ± 0.2[b]	372 ± 16[a]	7.6 ± 0.8[c]
G2 (30-30A-30A)	19.4 ± 0.2[b]	31.6 ± 3.4[b]	4.3 ± 0.6[b]	358 ± 13[a]	9.0 ± 1.4[c]
G3 (30A-30A-30A)	17.9 ± 1.2[b]	31.1 ± 2.3[b]	4.5 ± 0.2[b]	367 ± 22[a]	8.5 ± 1.6[c]
G4 (30A-30-30)	20.2 ± 0.1	35.6 ± 2.1	4.8 ± 0.3[b]	489 ± 11[b]	4.4 ± 0.1[b]
G5 (90)	20.1 ± 0.3	32.4 ± 2.6	4.6 ± 0.4[b]	483 ± 15[b]	4.8 ± 0.1[b]
	$P \leq 0.005$[a,b]	$P \leq 0.02$[a,b]	$P < 0.0001$[a,b]	$P < 0.0001$[a,b]	$P \leq 0.0002$[a,b]
		$P > 0.54$[b]			$P \leq 0.0008$[b,c]

[a, b; b; b, c]Levels of significance (P-values) are given for statistical comparisons between material properties labeled with the corresponding superscripts.

reduction in fatigue strength compared to G1 and unirradiated UHMWPEs. Furthermore, each annealing step appeared to decrease the slope of the *S–N* curve (Figure 5). Most likely, annealing treatments were responsible for the main decrease in fatigue resistance, since irradiation without further annealing has been demonstrated to slightly augment the fatigue strength of e-beam irradiated UHMWPEs.[9] Despite the registered drop in fatigue resistance upon annealing, sequentially crosslinked UHMWPE, G3, appeared to have long-term fatigue performance closer to that displayed by unirradiated UHMWPE specimens. As proposed elsewhere, there seems to be a direct relation between mechanical behavior and microstructure for highly crosslinked UHMWPEs.[8,9,35] Thus, irradiation results in crystal thickening, which, in turn, is responsible for an improvement of the fatigue life in long-term experiments. Also, a 2 h annealing has been reported to imply a decrease in lamellar thickness compared to as-irradiated UHMWPE, and, coherently, to demonstrate a reduced fatigue strength.[9] The negative impact on fatigue behavior of longer annealing steps (8 h) found in this study appear to be compensated to some extent introducing the sequential irradiation-annealing strategy.

The microstructural changes induced by irradiation caused substantial reductions in crack propagation resistance behavior, regardless of the stabilization strategy. FCP results showed two different regions in the log–log plots of crack growth rate, (da/dN), versus stress intensity factor range (ΔK) (Figure 6). The first region matched the slow crack growth regime, whereas the second one represented the intermediate crack growth or Paris equation regime ($da/dN = C\,(\Delta K)^m$; C and m are constants). A fatigue crack inception stress intensity range ($\Delta k_{\text{inception}}$) could be defined as the intersection of the first regime, nearly vertical, curve with the x-axis, at a value of $da/dN = 10^{-9}$ m/cycle. This approach gave the stress intensity threshold that must be overcome to initiate the propagation of a static crack and permitted comparison between materials. The second region fitted to a

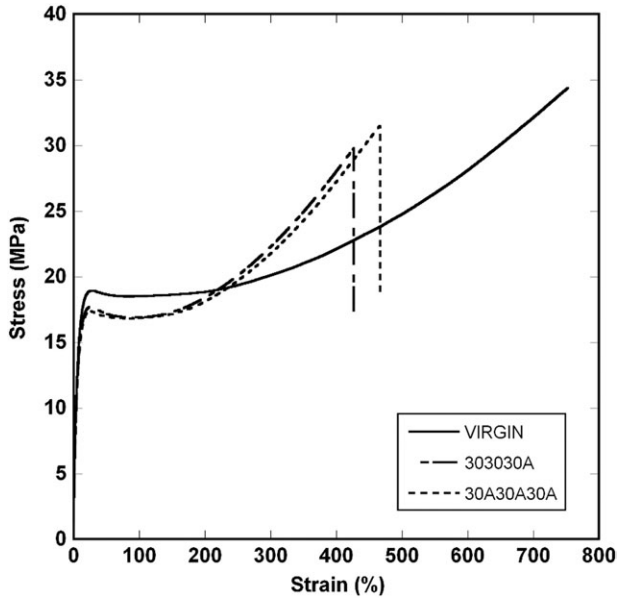

Figure 4. Typical engineering stress–strain curves obtained from uniaxial tensile tests for virgin, postirradiation annealed, and sequentially crosslinked UHMWPEs.

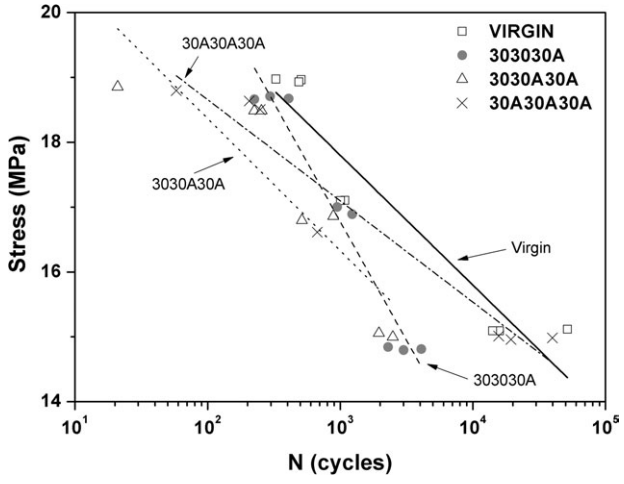

Figure 5. Stress–life curves for virgin, postirradiation annealed, and sequentially crosslinked UHMWPEs.

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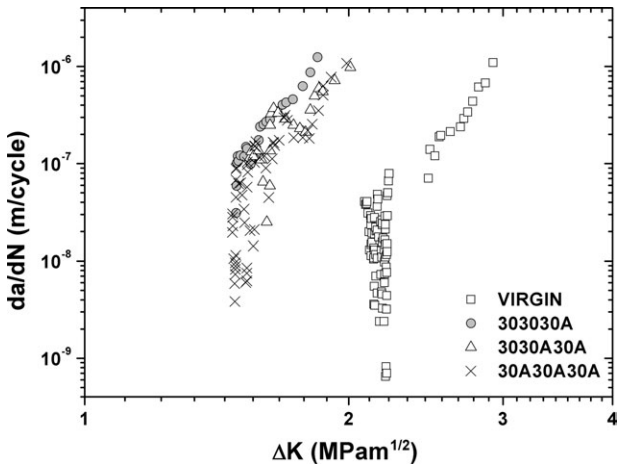

Figure 6. FCP curves corresponding to virgin, postirradiation annealed, and sequentially crosslinked UHMWPEs.

linear trend, whose slope, *m*, provided information about how fast the crack propagate once it started to grow. Fast-fracture regime was reached at the end of all the experiments. The unirradiated material presented the highest Δk at crack inception (2.2 ± 0.1 MPa/m$^{1/2}$), whereas the corresponding Δk of the crosslinked UHMWPEs (G1, G2, and G3) dropped to values close to 1.6 MPa/m$^{1/2}$ (Table III). This finding was not unexpected, as previous studies have confirmed remarkably drops in stress intensity factor at crack inception, $\Delta k_{inception}$, after irradiation.[8–11,35] Thus, the greater the radiation dose, the higher crosslink density and the lower the $\Delta k_{inception}$. The elevated crosslink density imparted by irradiation reduces the deformation modes of the amorphous region, and therefore cracks grow more easily in crosslinked UHMWPEs.[36] Conversely, no significant differences ($P > 0.05$) were found regarding the crack inception behavior among crosslinked UHMWPEs. It can be concluded that the annealing strategy, terminal or sequential, scarcely affected the crack propagation resistance, confirming crack inception is mainly governed by crosslink density in crosslinked UHMWPEs. Finally, sequentially crosslinked materials, G2 and G3, had less steep slopes (lower *m* coefficients) than unirradiated and postirradiation annealed UHMWPEs (Table III). This fact might indicate that lower stress levels are needed to reach similar crack growth rates.

Table III. Stress–Intensity Levels at Crack Inception (Mean ± SD) for Virgin, Postirradiation Annealed, and Sequentially Annealed UHMWPEs

Material	$\Delta K_{inception}$ (MPa/m$^{1/2}$)	*m*
Unirradiated	2.22 ± 0.06	12.1 ± 2.0
G1 (30-30-30A)	1.49 ± 0.03	12.9 ± 0.8
G2 (30-30A-30A)	1.58 ± 0.08	8.6 ± 0.3
G3 (30A-30A-30A)	1.51 ± 0.03	7.1 ± 1.5
B100A[a]	1.49 ± 0.06	12.9 ± 0.8

[a]Data reported in Ref. 9 corresponding to 100 kGy e-beam irradiated and annealed UHMWPE.

Table IV. Estimations of Toughness Behavior (Mean ± SD) from Uniaxial Tension and Impact Tests

Material	Work to fracture (MPa or MJ/m^3)	Impact toughness (kJ/m^2)
Unirradiated	209.7 ± 20.7[a]	100.9 ± 10.3[a]
G0 (30-30-30)	116.5 ± 4[b]	62.6 ± 4.7[b]
G1 (30-30-30A)	88.8 ± 6.4[b]	47.1 ± 2.9[b, c]
G2 (30-30A-30A)	93.3 ± 19.4[b]	47.8 ± 2.6[b, c]
G3 (30A-30A-30A)	94.1 ± 7.1[b]	47.9 ± 5.2[b, c]
G4 (30A-30-30)	114.8 ± 10.7[b]	N/A
G5 (90)	105.3 ± 12.8[b]	N/A
	$P < 0.0001$[a,b]	$P < 0.0001$[a,b]
		$P \leq 0.0015$[b,c]

N/A: Not available.
[a, b, b, c]Levels of significance (P-values) are given for statistical comparisons between material properties labeled with the corresponding superscripts.

Toughness Behavior

The radiation dose absorbed was the key parameter governing the fracture resistance of the various UHMWPE formulations. A significant decrease in impact toughness was confirmed for crosslinked UHMWPEs compared to the unirradiated polymer ($P < 0.0001$). Thus, impact toughness dropped about a 40% upon irradiation and a 50% after crosslinking and stabilization processes (Table IV). As mentioned before, the elevated crosslink density limits the ductility, and also its fracture resistance, as the crosslinked network prevents the polymer from reaching high deformations.[35] Crystallinity drops registered after performance of annealing treatments also negatively affected the fracture resistance of crosslinked UHMWPEs, although to a much lesser extent. However, the introduction of more than one annealing step had no further influence on impact toughness, as no significant differences could be detected among crosslinked and stabilized UHMWPEs regardless of the annealing strategy. Work to fracture results followed a trend similar to that of impact results. Unirradiated UHMWPE had the highest work to fracture ($P < 0.0001$), whereas crosslinked UHMWPE exhibited very low values, mostly due to the loss in ductility. The energy needed to fracture crosslinked materials decreased a little bit further as more annealing steps were introduced (Table IV). Finally, load displacement curves to fracture corresponding to compact tension specimens revealed a similar behavior, with unirradiated UHMWPE needing high loads and displacements to reach fracture, while crosslinked UHMWPEs had much lower values (Figure 7). Again, no significant differences were found among crosslinked UHMWPEs.

Commercially available sequentially crosslinked UHMWPE is claimed to have excellent oxidation resistance (no detectable free radicals) and superior mechanical properties compared to its postirradiation annealed predecessor.[23,24,37] It is worth mentioning that the UHMWPE resins used to produce these formulations are different. The manufacturer replaced the GUR 1050 UHMWPE resin employed to fabricate postirradiation annealed UHMWPE with GUR 1020 resin to produce the sequentially annealed formulation. The latter resin is a lower molecular weight powder, and UHMWPE materials produced from this

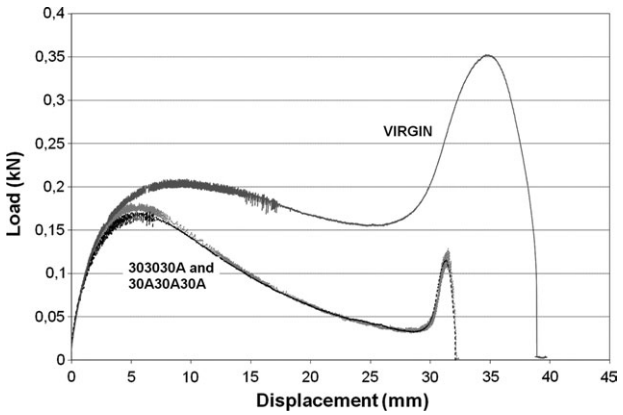

Figure 7. Load–displacement curves to fracture corresponding to compact tension specimens of virgin, postirradiation annealed, and sequentially crosslinked, G3, UHMWPEs.

resin have been reported to exhibit improved mechanical properties than those manufactured from GUR 1050.[38] So, it is unclear whether the improvement stems from the annealing strategy or the UHMWPE resin. Our study suggests that oxidation resistance seems to be superior for sequentially annealed UHMWPEs from a thermooxidation perspective. However, the mechanical improvement obtained after introduction of three sequential irradiation-annealing steps appears to be quite limited, at least when GUR 1050 resin is used as base material.

Obviously, the present study is not free of some limitations. First, thermooxidation parameters might not necessarily correlate with oxidation indices measured after shelf-aging or *in vivo* oxidation conditions. The orthopedic community generally relies on standard accelerated aging protocols to explore the oxidative stability of alternative polyethylenes and to categorize them. However, accelerated aging protocols have not always provided an exact correspondence to oxidation indices and regional distribution of oxidation maxima found in shelf aged implants or retrievals. Remelted polyethylenes represent an interesting example as these materials performed very well after accelerated aging,[39] but recent evidence suggests no complete oxidation resistance was achieved.[40] We chose to perform thermogravimetry since it provides a faster first screening regarding oxidative stability of the molten polymer. Second, cyclic stress–strain and long-term fatigue experiments are not intended to confirm the suitability of the studied polyethylene materials as acetabular liners or tibial inserts, or to predict an optimal clinical performance. These mechanical tests do not take into account complex load patterns (biaxial or triaxial stress states), and, conversely, the clinical performance of the artificial joint depends on a variety of patient, surgical, design and material factors.

CONCLUSIONS

This study provides evidence that the introduction of sequential irradiation-annealing processes may improve the resistance to oxidation as compared to postirradiation annealed UHMWPEs. The microstructural characterization of sequentially crosslinked UHMWPEs also confirmed crystal thickness and crystallinity contents similar to those of the unirradiated polymer, whereas the TG behavior suggested this material had the highest crosslink density. The anticipated improvement in mechanical properties, however, appears to be more limited, as the mechanical, crack propagation, and fracture resistance properties were generally comparable to those of postirradiation annealed and G2 (two sequential irradiation and annealing steps) UHMWPEs.

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