Regulated splicing of the fibronectin EDA exon is essential for proper skin wound healing and normal lifespan

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Fibronectins (FNs) are multifunctional high molecular weight glycoproteins present in the blood plasma and in the ECMs of tissues. The FN primary transcript undergoes alternative splicing in three regions generating up to 20 main different variants in humans. However, the precise role of the FN isoforms is poorly understood. One of the alternatively spliced exons is the extra domain A (EDA) or extra type III homology that is regulated spatially and temporally during development and aging. To study its in vivo function, we generated mice devoid of EDA exon-regulated splicing. Constitutive exon inclusion was obtained by optimizing the splice sites, whereas complete exclusion was obtained after in vivo CRE-loxP-mediated deletion of the exon. Homozygous mouse strains with complete exclusion or inclusion of the EDA exon were viable and developed normally, indicating that the alternative splicing at the EDA exon is not necessary during embryonic development. Conversely, mice without the EDA exon in the FN protein displayed abnormal skin wound healing, whereas mice having constitutive inclusion of the EDA exon showed a major decrease in the FN levels in all tissues. Moreover, both mutant mouse strains have a significantly shorter lifespan than the control mice, suggesting that EDA splicing regulation is necessary for efficient long-term maintenance of biological functions.

Introduction

Fibronectins (FNs)* are high molecular weight glycoproteins that are present on many cell surfaces, in extracellular fluids, connective tissues, and basement membranes. They interact with other ECM proteins and cellular ligands, such as glycosaminoglycans, collagen, fibrin, and integrins (Kornblihtt and Gutman, 1988; Hynes, 1990). FNs play key roles in the adhesive and migratory behavior of cells related to fundamental processes such as embryogenesis, malignancy, homeostasis, wound healing, host defense, and maintenance of tissue integrity (Hynes, 1990; ffrench-Constant, 1995; Kornblihtt et al., 1996). FN molecules are a mixture of several protein types that differ in both their primary structure and post-translational modifications. The amino acid sequence variations are the consequence of the alternative processing of a single primary transcript at three sites: extra domain B (EDB) or extra type III homology B (also called EIII-B or EDII), extra domain A (EDA [EIII-A or EDB]) and type III homologies connecting segment (IIICS; V region in rat). The alternative splicing process is cell type-, developmentally, and age regulated. The EDA and EDB are single exons coding for single type III repeats that are included or excluded from the FN mRNA by exon skipping (Kornblihtt et al., 1984; Gutman and Kornblihtt, 1987; Schwarzauer et al., 1987; Zardi et al., 1987). The IIICS region undergoes a more complex pattern of splicing which is species dependent (Schwarzauer et al., 1983). In humans, over 20 main different isoforms are generated after mRNA processing, whereas in rats and mice, 12 variants are produced (ffrench-Constant, 1995; Kornblihtt et al., 1996). Two major forms of FN exist, plasma (pFN) and normal lifespan...
and cellular (cFN), having similar but not identical polypeptides of ~220–240 kD. pFN is a soluble dimeric form that lacks both the EDA and EDB domains. It is synthesized by hepatocytes and secreted into the bloodstream. cFN is a dimeric or cross-linked multimeric form containing the EDA and EDB domains at variable proportions. It is made by fibroblasts, epithelial, and other cell types, and is deposited as fibrils in the ECM (Hynes, 1990; ffrench-Constant, 1995; Kornblith et al., 1996).

The molecular mechanisms involved in processing the FN transcript are well understood for both the EDA and EDB exons (Mardon et al., 1987; Lavigueur et al., 1993; Caputi et al., 1994; Du et al., 1997; Lim and Sharp, 1998; Muro et al., 1999). The mouse EDA exon is regulated in a similar manner to its human counterpart and optimization of the 5’ and 3’ splice junctions of the exon leads to constitutive splicing in NIH3T3 cells (Muro et al., 1998).

Despite the fact that both FN primary structure and FN alternative splicing were described almost two decades ago (Kornblith et al., 1983; Petersen et al., 1983; Schwarzbauer et al., 1983; Kornblith et al., 1984; Zardi et al., 1987), many aspects of the function of FN isoforms remain elusive. In vivo studies in chicken, rat, and Xenopus showed the increased inclusion of EDA and EDB exons in the FN mRNA derived from embryos during development (ffrench-Constant and Hynes, 1989; Oyama et al., 1989; Pagani et al., 1991; DeSimone et al., 1992). Once development is complete, the inclusion of EDA and EDB decreases in a wide range of tissues. In fact, EDA- and EDB- forms are the most abundant forms in adult humans and rats (Magnuson et al., 1991; Pagani et al., 1991). This exclusion is cell type specific and differs in extent between the two exons. Quantitative mRNA studies in these tissues showed that the EDB exon is excluded more often than the EDA exon from the FN mRNA (ffrench-Constant and Hynes, 1989; Pagani et al., 1991; Caputi et al., 1995). It has been postulated that the inclusion of the spliced regions could alter the conformation of the RGD sequence, which is the main cell-binding site, via modification of its neighboring modules (ffrench-Constant, 1995) or by the alteration of the global conformation of the FN molecule (Manabe et al., 1997). In both cases, the alteration might affect the strength of the central cell–binding domain interaction with integrins. Since integrins α5β1 and αvβ3 were recently described as the cellular receptors for the EDA segment, another possibility is that cell adhesion can be directly regulated by alternative splicing (Liao et al., 2002). Other functions proposed for the EDA segment are as follows: wound healing (Clark et al., 1983; ffrench-Constant et al., 1989); matrix assembly (Guan et al., 1990); dimer formation (Peters et al., 1990); secretion (Wang et al., 1991); cell adhesion (Xia and Culp, 1995); cell differentiation (Jarnagin et al., 1994); tissue injury and inflammation (Satoi et al., 1999; Okamura et al., 2001); and cell cycle progression and mitogenic signal transduction (Manabe et al., 1999). Because most of the above studies were done using in vitro and cell culture systems, the in vivo role of the EDA segment still remains obscure. The use of mouse models could provide the best approach to understand the in vivo function of the EDA exon. However, the in vivo study of the function of the different protein isoforms is complex due to the simultaneous presence of more than one protein form at a given specific developmental time and tissue. Gene targeting in mice has allowed the specific deletion of alternatively spliced exons in different protein systems such as α5 integrin, γ-aminobutyrate receptor, dopamine D2 receptor, FN, Pax6, and Stat3 (Gimond et al., 1998; Homanics et al., 1999; Usiello et al., 2000; Wang et al., 2000; Fukuda et al., 2002; Singh et al., 2002; Yoo et al., 2002). Thus, it represents the best approach to understand the in vivo function of the EDA exon.

Taking advantage of our detailed knowledge of the elements involved in EDA splicing regulation (Mardon et al., 1987; Caputi et al., 1994; Muro et al., 1998, 1999), we have designed a novel approach to study the in vivo function of protein isoforms coded by genes that undergo alternative splicing without modifying the coding sequence of the FN gene. Using gene targeting, we generated a mouse strain containing the EDA allele with optimized splice sites at both splicing junctions (EDAα– allele), with loxP sites located in the adjacent introns. By mating this mouse strain with a “deleter” Cre-recombinase expressing mouse, an EDA-null allele was obtained (EDA−/− allele). Here, we show that FN mRNA produced by homozygous EDA+/− mice contained constitutive inclusion of the EDA exon devoid of developmental and tissue-specific regulation. Mice lacking EDA splicing regulation (EDA+/− and EDA−/−) were viable and phenotypically similar to the wild-type mice. However, the mouse strain producing EDA−FN showed an abnormal cutaneous skin wound healing. On the other hand, mice having the EDA exon constitutively included in the FN mRNA showed normal wound healing but had a striking decrease in the levels of FN in most of the organs analyzed from adult mice. Furthermore, and most importantly, both mouse strains had a significantly shorter lifespan than EDA+/−/+ animals, suggesting that the presence of both isoforms was necessary for efficient long-term maintenance of biological functions.

Results
The study of the in vivo function of protein isoforms generated by alternatively spliced genes is generally limited by the fact that the different isoforms coexist in a temporal and spatial manner in the same tissue of the organisms. Therefore, the availability of animal models having a controlled expression of the different forms is crucial for these studies. We have previously reported that optimization of the 5’ and 3’ splicing sites of the mouse EDA exon from the FN gene leads to constitutive exon inclusion in NIH3T3 cells (Muro et al., 1998). These results coupled with the power of homologous recombination technology in mouse embryonic stem (ES) cells enabled us to study the biological significance of protein isoforms generated by alternative splicing (for a detailed description of the rationale see Online supplemental material available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1). Furthermore, to date, there is no report of optimization of splice sites leading to constitutive splicing in vivo of a naturally alternatively spliced exon.
In vivo function of the EDA exon of the FN gene

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Mouse strains lacking alternative splicing at the EDA exon develop normally and are fertile

To generate an FN allele devoid of alternative splicing in the EDA exon, the wild-type EDA exon in the FN gene was replaced with a “floxed” EDA exon having both the 5' and 3' optimized splice sites described in Fig. S1 (available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1). After electroporation of 129 Sv/J ES cells with the targeting construct (Fig. 1 A), G418-resistant clones were selected and analyzed by Southern blot hybridization. A second step of recombination by transient expression of the CRE-recombinase was performed to eliminate the Neo-TK cassette from the EDA-flanking intron. Southern blot analysis and sequencing of the PCR-amplified genomic EDA region confirmed the introduction of the optimized 3' and 5' splice sites and the loxP sites flanking the EDA exon in one of the FN alleles. Hereafter, this allele will be called the EDA⁺⁻ allele. The heterozygous EDA⁺⁻ ES cells were used to obtain male chimeras that transmitted the EDA⁺⁻ allele to the germline. Heterozygous EDA⁺⁻ mice were mated with CMV-CRE-transgenic mice to obtain the EDA-null allele (henceforth, called the EDA⁻⁻ allele) after in vivo deletion of the loxP-flanked EDA exon. Homozygous mice were obtained by interbreeding heterozygous mice, and screening was performed by Southern blot analysis (Fig. 1 B) or by PCR of tail DNA biopsies (not depicted).

The homozygous EDA⁺⁻ and EDA⁻⁻ mice after birth exhibited no apparent developmental abnormalities compared with wild-type or heterozygous littermates; they reached adulthood and were fertile. In addition, no differences were detected in the organs from homozygous EDA⁺⁻ and EDA⁻⁻ mice by gross histological and pathological analysis (unpublished data). The mutated alleles segregated at the expected Mendelian frequency, and litter size and survival rate of newborn mice (measured until weaning of the litter) were similar to that of EDA⁺⁺/wt animals. No differences were observed neither in the shape of the embryos nor in the number of somites of embryos derived from timed matings. The detailed data are shown in the Online supplemental material section.

EDA⁺⁻ mice lack alternative splicing of the EDA exon

The inclusion of the EDA exon within the FN mRNA is subjected to a tight tissue-specific regulation. The most dramatic example is seen in liver, where pFN is synthesized and the EDA exon is completely excluded from the mRNA in the adult (Tamkun and Hynes, 1983; Kornblith et al., 1984).

Radioactive RT-PCR was used to analyze most of the tissues and organs in all three genotypes (EDA⁺⁺/wt, EDA⁺⁻, and EDA⁻⁻). An example of the EDA splicing pattern in representative tissues such as liver, brain, and kidney is shown in Fig. 1 C. These results were confirmed by RNase protection analysis of the same samples (Fig. S2 available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1). We observed that the EDA⁺⁺ form was never present in the samples from EDA⁻⁻ mice and the EDA⁺⁺ product from EDA⁺⁻ mice was present in all tissues. Both RNase protection and RT-PCR analysis of the FN mRNAs showed that there is 99–100% inclusion of the EDA exon in all tissues from EDA⁺⁻ mice, whereas there is 100% EDA⁺⁺ exon exclusion in all tissues belonging to the EDA⁻⁻ mice. This confirmed that the rationale of genetic modification was successfully fulfilled in the real in vivo situation.

Normal FN-ECM formation in mouse embryonic fibroblasts (MEF) derived from mice lacking regulated splicing at the EDA exon

Next, we examined the expression of FN mRNA, protein isoforms and ECM formation in MEF prepared from EDA⁺⁺/wt, EDA⁺⁻, and EDA⁻⁻ animals. RT-PCR analysis confirmed the constitutive splicing-in of the EDA exon in the EDA⁺⁻ and its absence in the EDA⁻⁻ MEFs (Fig. 2 A). Analysis of protein extracts by Western blot using polyclonal anti-FN...
Regulated splicing of the EDA is dispensable for the ECM formation. (A) RT-PCR analysis of total RNA prepared from MEF from EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> embryos (13.5 d post coitus [p.c.]). (B and C) Western blot analysis of protein extracts prepared from the above-described MEF (20 and 100 µg for B and C, respectively) with anti-FN and anti-EDA antibodies, respectively. Coomassie blue staining showed equal loading. (D) MEF prepared from EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> embryos (13.5 d p.c.) were plated, fixed, and incubated with anti-EDA antibody (left column) or with anti-total FN antibody (middle column). The merged image is shown in the right column.

and monoclonal anti-EDA antibodies showed normal protein levels in both the EDA<sup>+/+</sup> and EDA<sup>−/−</sup> MEF (Fig. 2 B). The anti-EDA antibody detected the EDA<sup>+</sup> FN isoform only in EDA<sup>+/+</sup> and EDA<sup>wt/wt</sup> MEF (Fig. 2 C). Immunofluorescence analysis using a confocal microscope showed the presence of a normal FN-ECM in EDA<sup>+/+</sup> and EDA<sup>−/−</sup> MEF when compared with EDA<sup>wt/wt</sup> MEF (Fig. 2 D). The signal obtained with the anti-EDA antibody localized with that observed with the anti-FN antibody.

Ablation of regulated splicing in the EDA exon produced a decrease in FN levels in EDA<sup>+/+</sup> mice that did not correlate to changes neither in FN mRNA nor in the integrin levels.

Next, we examined whether the modifications in the natural alternative splicing pattern of FN had some consequences in other cellular proteins or FN itself. Consequently, protein extracts from different tissues of the different genotypes were

Figure 2. Regulated splicing of the EDA is dispensable for the ECM formation. (A) RT-PCR analysis of total RNA prepared from MEF from EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> embryos (13.5 d post coitus [p.c.]). (B) Western blot analysis of protein extracts prepared from the above-described MEF (20 and 100 µg for B and C, respectively) with anti-FN and anti-EDA antibodies, respectively. Coomassie blue staining showed equal loading. (D) MEF prepared from EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> embryos (13.5 d p.c.) were plated, fixed, and incubated with anti-EDA antibody (left column) or with anti-total FN antibody (middle column). The merged image is shown in the right column.

Figure 3. The decrease in FN levels in tissues of EDA<sup>+/+</sup> mice is not correlated to a reduction in integrin levels. (A and B) Western blot analysis of total protein extracts (50 µg) from brain, heart, lung, and liver from all genotypes (EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, EDA<sup>−/−</sup>, EDA<sup>+/−</sup>, EDA<sup>−/−</sup>, and EDA<sup>+/−</sup>) using anti-FN polyclonal antibody (A) and, after stripping the membrane, anti-β-tubulin mAb to verify for minor errors in protein load (B). Note that the migration of the EDA<sup>+</sup> FN is slightly slower than that of the EDA<sup>−</sup> FN. Coomassie blue staining of 5–17% gradient gels of the same protein extracts did not show changes in other proteins. (C) Plasma and serum protein samples from EDA<sup>+/+</sup>, EDA<sup>−/−</sup>, and EDA<sup>wt/wt</sup> mice (20 µg) were Coomassie blue stained (left), and a similar gel was blotted and incubated with anti-FN antibody (middle). For the anti-EDA antibody (right), 100 µg of protein extract were loaded. (D and E) Western blot analysis of α<sub>9</sub>- and β<sub>1</sub>-integrin levels in different tissues (liver, brain, and skin) using anti-α<sub>9</sub>- and β<sub>1</sub>-rabbit polyclonal antibodies. Anti-β-tubulin mAb was used to verify for minor errors in protein load.
prepared from 3-mo-old mice and analyzed by SDS-PAGE. No evident difference was observed at least at the sensitivity level of Coomassie blue staining of 5–17% gradient gels (unpublished data). On the other hand, a Western blot analysis using anti-FN polyclonal antibody showed a striking decrease in FN levels in most of the tissues analyzed from EDA+/+ mice (Fig. 3 A). This decrease was also evident in EDA+/+ and EDA−/− mice, both bearing one EDA+ allele. The observed differences were not due to a different protein load of the gels, as evidenced by the Coomassie blue staining and by the incubation of the same membranes with an anti–β-tubulin mAb (Fig. 3 B). A significant decrease in FN amounts was also observed in the plasma and serum samples prepared from EDA+/+ animals when compared with those from EDA+/+ mice (Fig. 3 C). Western blot using an anti-EDA antibody confirmed the presence of EDA containing FN in the EDA+/+ plasma and its complete absence in the EDA−/− sample (Fig. 3 C).

It was considered important to analyze if the FN level differences observed in the different tissues were associated to changes in the levels of the specific EDA receptor α9β1-integrin (Liao et al., 2002). We observed no differences in α9- and β1-integrin levels among EDA+/+, EDA+/−, and EDA−/− mice by Western blot analysis in liver and brain (Fig. 3 D). However, it remains to be seen if the localization pattern of the integrin receptors or their signaling have been affected in the mutant mice.

On the other hand, to determine whether the observed decrease in FN protein levels in the EDA+/+ mice was correlated to a decrease in the mRNA levels, we performed Northern blot analysis of total RNA isolated from the tissues having the higher decrease in FN levels (brain and heart) from 3-mo-old male mice. No significant variations were

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**Figure 4.** Expression of FN in different tissues at different ages. (A) Protein extracts from liver, heart, lung, and brain were prepared from EDA+/+, EDA+/−, and EDA−/− mice of different ages (13.5 d p.c., 1, 3, and 14 mo old) and analyzed by Western blot using an anti-FN antibody. (B) The histograms represent the relative amount of FN present in EDA+/− mice relative to that present in EDA+/+ mice (considered as 100%) for each tissue and each time point.

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**Figure 5.** Reepithelization is observed 5 d after full thickness cutaneous wounding. Full thickness cutaneous wounds of control (EDA+/+, n = 6) and mutant mice (EDA+/− and EDA−/−, n = 6 and 7, respectively) mice were analyzed at 5 d after wounding by hematoxylin-eosin staining. Representative sections are shown. The black arrowheads indicate the wound edges. Epithelium, granulation tissue, dermis, newly formed epithelium, and eschar are indicated (“e,” “g,” “d,” “re,” and “sc,” respectively). The dotted rectangles in the top panels indicate the magnified area showed in the bottom panels. The experiment was repeated twice with similar results. Edematous granulation tissue is observed in the EDA−/− wounds.
Figure 6. EDA−/− skin wounds at day 7 after wounding show abnormal healing and increased number of BrdU-positive cells. (A) Full thickness cutaneous wounds of control (EDA<sup>wt/wt</sup>, n = 8) and mutant mice (EDA<sup>+/−</sup> and EDA<sup>−/−</sup>, n = 8 and 9, respectively) mice were analyzed at 7 d after wounding. Representative sections are shown. The black arrowheads indicate the wound edges. Abbreviations are as described in Fig. 5. The ulcerated epithelium observed in the EDA<sup>−/−</sup> wound is indicated as “ue.” The experiment was repeated three times with similar results. Wound sections of three independent experiments (eight mice per genotype in each experiments) were microscopically scored for the presence of ulcerative processes at day 7 after wounding. The fraction of ulcerated wounds of the EDA<sup>−/−</sup> mice was statistically significant different from that of EDA<sup>wt/wt</sup> and EDA<sup>+/−</sup> mice (63 and 22%, for EDA<sup>−/−</sup> and EDA<sup>wt/wt</sup>, respectively; P < 0.0002 for EDA<sup>−/−</sup> vs. EDA<sup>wt/wt</sup> by both...
observed in RNA samples from the brain and heart of EDA^{+/+} mice (Fig. S3 available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1).

Embryos and prepubertal EDA^{+/+} mice have normal FN levels
To determine whether the mechanisms that produced the FN decrease were developmentally regulated, we performed a Western blot analysis of samples prepared from mice of different ages. The decrease in FN levels was not present during embryonic development. It started after birth in most of the tissues, and arrived to a plateau in 2-mo-old mice (Fig. 4). In some tissues, the FN levels in EDA^{+/+} mice were \( \sim 10\%-20\% \) of the amount found in EDA_{wt}/wt mice (heart, blood, brain, kidney, and diaphragm), whereas the liver showed a lesser decrease in FN levels in EDA^{+/+} mice, arriving to 60\%-70\% of the amount present in EDA_{wt}/wt mice.

In an attempt to determine alterations in the FN-degradation mechanisms, we performed gelatin zymography and observed no variations in plasma and other tissues (brain and heart) in the levels of matrix metalloproteinases (MMPs), enzymes known to degrade FN (Sternlicht and Werb, 2001). Moreover, no differences were observed in the degradation of in vitro translated recombinant fragments of FN with or without the EDA exon that were incubated with tissue extracts (unpublished data). Western blot analysis of tissue extracts using a polyclonal anti-FN antibody did not reveal the presence of extra bands or any smaller degradation product of EDA^{+/+} tissues, neither with a 6\% nor a 12\% SDS-PAGE gel (unpublished data). Additionally, we were unable to reproduce in vitro the reduction in FN levels after culturing heart fibroblasts of adult mice (from 2-, 4-, and 6-mo-old animals), as the cell extracts from EDA^{+/+} mice showed similar levels of FN to those of the EDA_{wt}/wt fibroblasts (unpublished data).

These experiments showed that the striking reduction in FN protein levels in EDA^{+/+} mice occurred after birth. The decrease in FN was not related to the mRNA levels present in each tissue or to a detectable EDA\(^{-}\)-dependent degradation, suggesting that alternative mechanisms should be responsible for the decrease of tissue FN in the EDA^{+/+} animals.

EDA-null mice have abnormal skin wound healing
The FN present in the skin of wild-type animals does not contain the EDA exon. However, the FN-EDA\(^{+}\) form is thought to participate actively in the reepithelialization process (Clark et al., 1983; French-Constant et al., 1989). To explore the role of the EDA exon in the wound healing process, we performed full thickness excision skin wounds in EDA_{wt}/wt, EDA\(^{-}\), and EDA^{+/+} mice. The wound healing process was analyzed at 1, 3, 5, 7, 10, and 14 d after wounding. Hematoxylin-eosin staining of tissues from full thickness wounds during the early phase of reepithelialization (days 1 and 3) did not show any differences in reepithelialization and neovascularization (unpublished data). During late phase (days 5 and 7), the reepithelialization and formation of granulation tissue were indistinguishable between EDA^{+/+} and EDA_{wt}/wt mice (Figs. 5 and 6). However, the abnormal healing process was observed in the EDA^{−} mice. In fact, at day 5, the newly formed epidermis completely covered the wound area in the EDA^{−} mice but the border between the new epidermis and the granulation tissue was not sharply defined as in the control or EDA^{+/+} mice. Moreover, the granulation tissue of the EDA^{−} wounds showed less compact cells with edematous-like areas below the interface between the epidermis and the derma granulomatous reaction (Fig. 5, bottom).

At day 7, the EDA^{−} wounds were clearly different from those from EDA_{wt}/wt and EDA^{+/+} mice (Fig. 6 A, bottom). In fact, the EDA^{−} wounds showed ulcerative processes at the newly formed epidermal region level, resulting in a delay in the reepithelialization. Infiltration of inflammatory cells such as polymorphonuclear leukocytes and macrophages were detected by microscopic analysis of the edematous-like and ulcerative regions, and a higher cell proliferative activity was observed by in vivo BrdU labeling in these regions (Fig. 6 B, bottom right). On the contrary, neither EDA_{wt}/wt nor EDA^{+/+} wounds presented a high number of BrdU-positive cells in the area below the newly formed epidermal region (Fig. 6 B, bottom left and center). The status of the basal lamina in the EDA^{−} skin was similar to that of the EDA_{wt}/wt, as seen by both the Azan-Mallory coloration and the specific staining of laminin in the same sections using a rabbit polyclonal antilaminin antibody. Cell death and apoptosis-specific analysis were performed by TUNEL and caspase-3 assays of the wound sections, respectively, and showed no significant differences in the number of positive cells between the different genotypes (Fig. S4 available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1). Scoring of the skin healing process of three independent experiments (eight 3-mo-old mice per genotype in each experiment) for the presence of ulceration in the wounds showed that EDA^{−} mice developed them 2.9 times more frequently than the EDA_{wt}/wt mice (63\% of EDA^{−} vs. 22\% of EDA_{wt}/wt, \( P < 0.0002 \) by both the Fisher Exact test and the \( \chi^2 \) test) at d 7 after wounding.

A limited number of 11-mo-old mice (five EDA_{wt}/wt and six EDA^{−} mice) that were subjected to the same wound healing protocol showed an even higher proportion in the frequency of wound ulceration in the EDA^{−} mice (83.3\% vs. 20\%, EDA^{−} and EDA_{wt}/wt; respectively, \( P \leq 0.01 \) by both the Fisher Exact test and the \( \chi^2 \) test). To determine whether integrin \( \alpha_\beta_1 \) levels were changed, we performed a Western blot analysis of untreated skin samples and observed similar levels of integrin \( \alpha_\beta_1 \) among all three genotypes (Fig. 3 E).

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Constitutive exclusion or inclusion of the EDA exon within the FN mRNA shortens the lifespan

The in vivo studies performed on rats show a critical decrease in EDA+/EDA− ratio after birth, with an additional gradual decrease of the ratio in some tissues that is correlated with the age of the animal (Magnuson et al., 1991; Pagani et al., 1991). We addressed the question if in EDA+/+ and EDA−/− mice, having no regulation of EDA splicing, the presence or absence of the EDA exon could have an effect on their lifespan. For this purpose, we performed a long-term experiment using 139 mice (EDAwt/wt, n = 39; EDA+/+, n = 47; and EDA−/− n = 53) that were caged separately throughout the experiment and the time of animal death was recorded. The Kaplan-Meier graphical representation of the data is shown in Fig. 7. It can be seen that up to 15-mo-old mice from all the genotypes did not show any difference in survival. However, from thereon, there was a decline in the survival of EDA+/+ and EDA−/− mice. At 30 mo, only seven and eight EDA+/+ and EDA−/− mice were alive, (10 and 12.5% survival, respectively), whereas 21 EDAwt/wt mice survived (53.8% survival). Log-rank test statistical analysis indicated that both EDA+/+ and EDA−/− mice lifespan curves were statistically significant when compared with EDAwt/wt mice with P ≤ 0.0005. Postmortem external examination showed that 20–25% of the EDA−/− mice had different skin hyperproliferative and spontaneous ulcerative lesions on their mid-dorsal skin. In case of the EDA+/+ and EDA−/− mice, <5% had any kind of skin lesions. Autopsies were performed on naturally deceased mice of all three genotypes. Liver tumors, intestinal obstruction, prostatic necrosis, and pulmonary edema were the most frequently observed features, but there was no significant difference in these pathologies of the internal organs between the three genotypes.

Discussion

The first evidence of the in vivo importance of FN was demonstrated by R. Hynes’s group who showed that FN-null mice die during embryonic development (George et al., 1993). The early embryo lethality makes it difficult to study the biological functions of FN in the in vivo models. Therefore, one of the best methodologies to study the function of FN is by selective modification of the gene. Along with this line, Sakai et al. (2001) described a mouse model lacking pFN which showed increased neuronal apoptosis and larger infarction areas after transient focal cerebral ischemia. In an attempt to elucidate the role of the EDB exon, mice lacking this exon were also produced, but showed no obvious abnormalities in vivo (Fukuda et al., 2002).

We have investigated the in vivo role of the EDA exon by constructing mouse strains unable to carry out EDA exon alternative splicing. The EDA+/+ strain showing constitutive expression of the EDA exon in all tissues was obtained without any modification of the coding sequences and gene structure. This novel approach avoided the difficulties found in a previous attempt to produce animals devoid of alternative splicing of the EDB exon in the FN gene where the modification of the gene structure through the introduction of a selection cassette within the introns produced an FN-null allele (Georges-Labouesse et al., 1996). A similar phenomenon has recently been observed in other genes (Lewandoski, 2001). None of these negative effects was observed in our mutant mice, indicating that the insertion of loxP sites within the flanking introns did not modify the normal pre-mRNA transcription and processing.

Moreover, we observed no obvious embryonic mortality nor malformation in any of the homozygous mutant mice analyzed (EDA+/+ and EDA−/− animals), suggesting that the alternative splicing regulation of the EDA exon is not essential for the developmental processes or that it could be compensated by other gene products. The absence of any obvious developmental abnormality in EDA+/+ and EDA−/− mice suggests that the information and postulated roles for the EDA exon obtained by using recombinant FN fragments (Introduction) cannot be extended automatically to the in vivo functions where more complex mechanisms are acting.

EDA+/+ mice have decreased levels of FN in all tissues

The observation of decreased levels of FN in adult EDA+/+ mice, but not in embryos and very young mice (<1 mo old) suggests that different control mechanisms of FN levels are acting before and after puberty. This decrease could be due to one of the following explanations or a combination of them: (1) a decrease in FN mRNA levels; (2) a decrease in the secretion rate of FN; or (3) an increase in extracellular degradation of FN.

A series of experiments have been performed to test which of these mechanisms, if any, was acting in the EDA+/+ mice. Northern blot experiments showed that the mRNA levels were not directly related to FN protein levels (Figs. 3 and 4 and Fig. S3). Protease levels were similar to those of EDAwt/wt, no additional FN degradation products were seen in EDA+/+ tissue extracts, and we were unable to see a decrease of FN levels in in vitro cultured cells derived from adult organs of EDA+/+ mice. The failure to identify the precise cause of the low FN-EDA+ levels suggests that an atypical mechanism might be acting. One possibility is that a specific intracellular trafficking mechanism is able to modulate the levels of EDA+ FN forms in adult animals. This mechanism was suggested for polarized secretion of FN forms in airway

![Image](https://example.com/image.png)

Figure 7. The lack of alternative splicing of the EDA exon reduces the lifespan of mutant mice. Both EDA+/+ and EDA−/− mice showed a reduced lifespan during the 30-mo period of analysis. The graphic shows a Kaplan-Meier representation of the survival versus time of 39, 45, and 53 EDAwt/wt, EDA+/+, and EDA−/− mice, respectively, that were housed in individual cages throughout the study. The results were analyzed with the Log-rank test and both the EDA+/+ and the EDA−/− curves were statistically significant when compared with the EDAwt/wt mice (indicated by asterisks, P ≤ 0.0005).
epithelial cells (Wang et al., 1991). It is interesting to note that Schwarzbauer et al (1989) have shown that the IIICS-0/IIICS-0 dimers were not secreted. It is possible that a similar control in the secretion mechanism could also be present for the EDA-1 subunits.

However, none of these effects was obvious in cultured fibroblasts from embryos and adult animals, and in the early stages of development of mice. In fact, the decrease in the amount of FN was evident only after 2–3 mo of age, which is after sexual development. Alternatively, a down-regulation of specific tissue inhibitors of MMPs in adult animals would have as its consequence an increase in activity of specific MMPs, a subtle change that direct gelatin zymography analysis will not detect.

**Cutaneous wound healing is abnormal in EDA-/- mice**

Initially, up to 3 d after wounding, there were no differences in the healing process between the three genotypes (EDAwt/wt, EDA-/-, and EDA+/+). This seems to be consistent with the fact that in the first days after wounding, deposited FN is mainly derived from plasma (Clark et al., 1983; ffrench-Constant et al., 1989). It was shown that normal wound healing could be helped by the presence of cFN (ffrench-Constant et al., 1989; Sakai et al., 2001). In fact, in mice devoid of pFN, normal wound healing was reported (Sakai et al., 2001). cFN is produced in situ 3 d after wounding (Clark et al., 1983; ffrench-Constant et al., 1989; Sakai et al., 2001) and a thin layer of cFN deposited on the surface of the wound immediately after injury in pFN-null mice (Sakai et al., 2001). Expression of cFN by the cells found at the base and edges of the wound just beneath the epidermis was observed, and the presence of cFN preceded the migration of epithelial cells that eventually covered the wound (ffrench-Constant et al., 1989). Previous studies have also shown that FN appears in the provisional matrix beneath the migrating epidermis coordinating with the FN-receptor expression on migrating epidermal cells (Clark et al., 1982, 1983; Grinnell, 1984). These data correlate perfectly with our results showing that after that period, despite the formation of granulation tissue and neovascularization, we observed abnormal reepithelialization in EDA-/- mice. The granulation tissue of the EDA-/- wounds displayed edematous regions that occur before the ulcerative process. The subsequent epidermal ulceration was accompanied by a proliferative stimulus with the influx of polymorphonuclear leukocytes and macrophages, and a delay in the healing of the wounds. Possible reasons for these phenomena could be related to defects in the basal lamina, a change in integrin receptor levels, and/or a defect in the skin keratinocytes, fibroblasts, or immune system. The basal lamina looked similar in EDA-/- mice compared with EDAwt/wt mice by both Azan-Mallory coloration and the specific immunostaining of laminin. However, we cannot rule out minor differences in one of the other specific proteins that are present in the basal lamina. Moreover, similar levels of α5β1 integrin, one of the EDA receptors (Liao et al., 2002), were observed in skin of the three genotypes. Preliminary attempts to define differences between the EDAwt/wt and EDA-/- genotypes in immune system cells (T/B lymphocytes ratio and delayed hypersensitivity) and in the characteristics of fibroblasts (in vitro cell migration, cell adhesion, and duplication time) failed to show significant differences. However, it remains to be seen if the localization pattern of the integrin receptors or their signaling have been affected in the mutant mice.

The increase in ulceration in the EDA-/- wounds could be the result of a higher rate of cell death in the skin of the mice during wound closure. This does not seem to be the case as the number of positive cells between the EDAwt/wt, EDA-/-, and EDA+/+ wounds in the granulation tissue were similar in the area devoid of ulceration by both TUNEL and caspase-3 assay.

On the basis of these considerations and the data described in Results, we conclude that FN containing the EDA segment plays an essential role in the organization of the granulation tissue and probably in epidermal cell migration, either directly or by interaction with other ECM components.

**EDA+/+ and EDA-/- mouse strains have a shorter lifespan**

It has been shown that deficiency of SH2-containing inositol-5-phosphatase, lysosomal acid lipase, and DNA topoisomerase IIIB had no consequences in embryonic development, but shortens the lifespan of the animals (Helgason et al., 1998; Du et al., 2001; Kwan and Wang, 2001). The shortened lifespan could be due to the changes in biological processes, which require the presence of the investigated protein during aging.

In the case of the FN mice, we showed that the absence of regulation of the EDA exon affects the lifespan of mutant animals. In normal animals, a dramatic change in the EDA+/EDA- ratio is observed after birth and a gradual decrease of EDA inclusion is observed in the tissues that increases with the age of the animal (Magnuson et al., 1991; Pagani et al., 1991). Alternative splicing of FN is regulated at both the extracellular (ffrench-Constant, 1995; Kornblitt et al., 1996; Boyle et al., 2000) and the molecular levels (Mardon et al., 1987; Lavigueur et al., 1993; Caputi et al., 1994; Muro et al., 1999). The mutant EDA+/+ and EDA-/- mice have no regulation of splicing of the EDA exon. It was possible that some biological processes that required specific EDA+/EDA- ratios of FN were altered because of the absence of regulation of EDA splicing. In fact, this was shown in this paper for the specific case of cutaneous wound healing in the EDA-/- mice. Furthermore, the low healing efficiency was exacerbated in older mice. In a similar fashion, other maintenance and tissue repair mechanisms could be affected, and the cumulative lower efficiency may result in a decrease in the lifespan of mutant animals. In the specific case of the EDA+/+ mice, due to the dramatic reduction in FN levels in most organs, it remains unclear whether the shortened lifespan seen in EDA+/+ mice was due to a change in FN quantity and/or quality. Although no significant differences have been reported on the lifespan of c129 and C57BL/6 (Smith et al., 1973; Goodrick, 1975; Kunsty and Leuenberger, 1975), we cannot formally rule out that the differences in chromosome background at the FN EDA locus (C57BL/6 for the EDAwt/wt and c129 with one or two loxP sites for EDA-/- and EDA+/+, respectively) could be responsible for the observed differences in lifespan.
In summary, we generated mice devoid of regulated splicing at the EDA exon of the FN gene. Surprisingly, both mouse strains showed that the alternative splicing at the EDA exon was dispensable during embryonic development, but it was necessary for a normal lifespan. Moreover, mice unable to incorporate the EDA exon in the FN protein displayed abnormal skin wound healing. On the other hand, mice having constitutive inclusion of the EDA exon showed an important decrease in the FN levels in all tissues.

The results presented here are important for three reasons: first, we presented a novel method to study the function of protein isoforms; second, we added valuable information regarding the function of the FN isoforms; and third, we showed that changes in the regulation of splicing are responsible for subtle changes in the phenotype that may be present, but undetected, in human pathologies.

Materials and methods
Mice
C57BL/6 mice were bred at the International Centre for Genetic Engineering and Biotechnology animal house or purchased from Harlan, Italy. All mice were housed in rooms at 25 °C with 12-h light/dark cycles. Mice were fed with standard food and water ad libitum. In all experiments, mice were males and age-matched, except in the aging experiment in which both males and females were used. Blood was drawn by puncture of the internal maxillary artery.

Generation of homozygous EDA mutant mice
Cloning of the murine genomic EDA region and targeting construct. A 13-kbp genomic clone containing the EDA exon region of the murine FN gene was isolated from a 129/Sv λ Dasmil library (a gift of G. Friedrich, Fred Hutchinson Cancer Research Center, Seattle, WA). The exon–intron structure and restriction map of this fragment was determined by restriction enzyme mapping followed by Southern blot analysis using individual exons as probes, which were PCR amplified from the human FN cDNA. Both the 5′ and 3′ splice sites of the EDA exon were modified by PCR-directed mutagenesis as described previously (Muro et al., 1998). In brief, the 5′ splice site was mutated to the 5′ consensus (5′/CAGgtatt3′) and the 3′ splice site was replaced by that of the constitutively spliced-in second exon of the apolipoprotein AI gene, which matches exactly the 3′ consensus (see Online supplemental material).

Our targeting construct consisted of 11.5 kbp of mouse genomic DNA. Neoymycin (Neo) and dihydrothiaxin (DTA) genes were used as positive and negative selection markers, respectively, for the first round of recombination. Three loxP sites were included in the targeting construct flanking the Neo-Thymidine Kinase (TK) cassette and the EDA exon.

The targeting construct was electroporated into ES cells, and 350 G418-resistant clones were genotyped by Southern blot hybridization of NcoI–SphI fragments using an internal probe (Fig. 1, A and B). F1 heterozygous offsprings were mated with C57BL/6 mice to obtain F2 generation, which were then mated to each other to produce EDA+/− mice (inclusion of the modified EDA exon in both the FN alleles). F1 EDA+/− mice were mated with a CRE-recombinase transgenic mice (on a C57BL/6 background) to obtain EDA+/− progeny (F2) that were subsequently mated to obtain EDA−/− animals.

Preparation of RNA from adult mouse and embryonic tissues
Total cellular RNA from various tissues was prepared by the guanidinium thiocyanate method (Chomczynski and Sacchi, 1987), quantitated by measuring absorbance at 260 and 280 nm, and RNA integrity was confirmed by running the samples on 1% agarose formaldehyde gels. For embryonic RNA samples, embryos were dissected under a stereomicroscope (model SMZ-10; Nikon) and tissues were pooled from five to six embryos (13.5 d p.c.). RNA was prepared using RNAZOL TM B (Biotex Labs) according to the manufacturer’s instructions.

mRNA analysis by Northern blot and radioactive RT-PCR
10–20 μg of total RNA from different tissues was denatured and run on a 1.2% agarose formaldehyde gel, blotted onto a nylon membrane (Hybond-N; Amersham Biosciences), and stained with a methylene blue solution to verify the efficiency of the transfer. The membrane was prehybridized in ExpressHyb™ hybridization solution (CLONTECH Laboratories, Inc.) and hybridized with the radioactive-labeled probe. It was exposed overnight using BIOMAX MS films (Kodak), and quantification of the radioactive signal was done with a phosphorimager (Canberra-Packard). FN cDNAs were analyzed as described in the Materials and methods section of the online supplemental material.

Preparation of protein extracts
Adult mouse tissues were surgically removed, homogenized in lysis buffer (150 mM NaCl, 50 mM Tris-HCl, pH 8, 1% wt/vol NP-40, and 2× complete protease inhibitor cocktail [Boehringer] with a polytron [Ultra-Turrax T8; Boehringer]), and analyzed by SDS-PAGE after protein determination using the Bradford assay (protein assay kit; Bio-Rad Laboratories). In the case of embryos, tissues from five to six embryos per genotype were surgically removed with the help of a stereomicroscope and pooled. The protein extracts were prepared by sonication in 150 μl of lysis buffer.

Western blot analysis
50 μg protein extracts were run on a 6%, 12%, and a gradient of 5–17% SDS-PAGE gels. The proteins were blotted onto a nitrocellulose membrane, incubated with the first and secondary antibodies, and the reaction was developed using the ECL system (Amersham Biosciences). The following primary antibodies were used: anti–human FN goat polyclonal antibody (Sigma-Aldrich), anti–human EDA mouse mAb 3E2 (Sigma-Aldrich), anti α-α-integrin rabbit polyclonal (a gift from D. Sheppard, University of San Francisco, San Francisco, CA), anti–β1-integrin rabbit polyclonal (Santa Cruz Biotechnology, Inc.), and the anti–β-tubulin mAb E7 (Developmental Studies Hybridoma Bank, University of Iowa) antibodies. Densitometry analysis was performed by using the Versa-Doc Analyzer (Bio-Rad Laboratories) with Quantity One software (Version 4; Bio-Rad Laboratories).

Immunofluorescence studies of MEF
Embryonic fibroblasts (13.5 d) from EDAwt/−, EDA−/−, and EDA−/− mice were cultured on coverslips for 24 h. Cells were washed and fixed with 4% vol/vol PFA in PBS, permeabilized with 0.5% vol/vol Triton X-100 in PBS, and washed. Cells were blocked with a mixture of anti–human FN and parallel with goat normal serum or with the mAb 3E2, and incubated with fluorescein-labeled rabbit anti-goat IgG (DakoCytomation) or with rhodamine-labeled rabbit anti-mouse IgG (DakoCytomation), respectively. Negative controls were performed with cells incubated directly with the fluorescent antibody. The analysis of the immunofluorescence was done by using a confocal laser scanning microscope (model Axiovert 100 M; Carl Zeiss Microimaging, Inc.).

Wound healing
Wounding and wound closure measurements. Full thickness skin wounds were done in 2–3-mo-old male mice (n = 12 per genotype) in two independent experiments. Genotypes were blind to the researcher and observer. In brief, animals previously anesthetized with Avertin, were shaved, swabbed with 70% ethanol and two circular, full thickness excisional wounds (5-mm diam, separated by 10 mm) were created on their backs, by excising the skin and panniculus carnosus. The mice were housed individually. For macroscopic analysis, mice were monitored daily and wounds were photographed and measured at each time point (0, 3, 5, 7, 10, and 14 d) to analyze the healing process.

Preparation, analysis, and histology of wound tissues. For histology of wounds, six 3-mo-old male mice from each genotype were killed at each time point (0, 3, 5, 7, 10, and 14 d), and their wounds were harvested by complete excision including the 3 mm of the epithelial margins. Wounds were immediately fixed in 2% formaldehyde in PBS containing 0.02% NP-
40 at 4°C for 3 h. They were placed in 2% formaldehyde in PBS, left overnight in the cold, and paraffin embedded. 3–5-mm sections from the middle of the wounds were stained with hematoxylin-eosin and analyzed by light microscopy. The analysis of the wound tissue at day 7 was performed in groups of eight mice per genotype in three independent experiments showing similar results. For basa lamina immunohistochemistry of the wounds, we used the polyclonal antilaminin antibody from Biogenex as described by the manufacturers.

**BrdU incorporation.** To allow the analysis of the cell proliferation in the wounds, the nucleoside analogue BrdU (250 mg/kg of body weight) was injected i.p. 2 h before the animals were killed (eight animals per each genotype, day 7 after wounding). Samples were prepared as described in the previous paragraph. Incorporation of i.p.-injected BrdU into DNA of replicating cells was analyzed using anti-BrdU mAbs conjugated with alkaline phosphatase as described by the manufacturer (BrdU-labeling and detection kit II; Roche). The day 7 time point with BrdU labeling was also performed with six 11-mo-old male mice per genotype. Representative data are shown in the figures.

**Statistical methods**

Data analyses were performed with the use of the software package GraphPad PRISM, version 2.0. Data were summarized with the mean as a measure of central tendency and the standard error as a measure of dispersion. The differences in proportions between the wild-type and mutant mice were assessed by using the χ² and the Fisher Exact test. A p-value of 0.05 was chosen as the limit of statistical significance. Log-rank test statistical analysis was used to evaluate the survival curves.

**Online supplemental material**

The online supplemental material section contains the rationale used for the optimization of splice sites and RNAse protection analysis of liver samples from control and mutant mice (see Figs. S1 and S2, respectively). The experimental analysis of allele frequencies, offspring, and embryos is also described. Northern blot analysis and TUNEL and caspase-3 analysis of skin wounds are shown in Figs. S3 and S4, respectively. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200212079/DC1.

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Xia, P., and L.A. Culp. 1995. Adhesion activity in fibronectin’s alternatively spliced domain EDa (EIIIA): complementarity to plasma fibronectin and plasma fibronec-
Supplemental results and discussion

Rationale for the optimization of the splice sites

We demonstrated previously that the mEDA3’SS(A) + 5’SScons plasmid, which contains a hybrid FN– α-globin mini-gene with optimized 5’ and 3’ EDA splicing sites, generates mRNA with constitutively included-in EDA exon (Muro et al., 1998). In this plasmid, the natural 5’ splice site (CAG:GTAT) was mutated toward the 5’ splice site consensus sequence (CAG:GTAAGT) to improve its base complementarity to U1 RNA (Fig. S1 C). In the same way, the 3’ splicing site was replaced by that of the constitutively spliced-in second exon of the ApoA1 gene, resembling the 3’ splice site consensus sequence. To obtain constitutive exclusion of the FN-EDA exon from the mRNA, we constructed the mEDA sequence. To obtain constitutive exclusion of the FN-EDA exon, we constructed the mEDA plasmid that had a deletion of 715 bp comprising the entire EDA exon (270 bp) and a small region of the flanking introns.

As there is tissue-specific regulation of the FN-EDA splicing pattern in vivo (Hynes, 1990; ffrench-Constant, 1995; Kornblihtt et al., 1996), both constructs were tested for efficiency of splicing by transient transfection in cell lines derived from different mouse organs having different EDA inclusion rate (unpublished data). Constitutive inclusion of the EDA exon was obtained with the mEDA3’SS(A) + 5’SScons plasmid in all cell types tested. As expected, the mEDAΔEDA plasmid produced only the EDA minus form in all cell types tested indicating that deletion of the EDA exon from the pre-mRNA did not alter the mRNA processing.

These results confirmed that both the optimization of the splicing sites of the EDA exon and its deletion lead to constitutive splicing independently of the cell type tested and opened the possibility of in vivo altering the alternative splicing pattern of the FN gene, and probably of other genes.

RNase protection analysis

We tested, by RNase protection analysis, the effectiveness of the EDA splicing site modifications in adult liver of the FN transcripts in all the genotypes (EDAwt/wt, EDA+/+, EDA−/−, EDA+/−, EDA−/wt, and EDA+/wt).

The RNase protection method is extremely sensitive and used to determine, in a quantitative manner, specific RNAs of the tissue under analysis without any further amplification (Sambrook et al., 1989). A 420-bp sequence-specific hybridization 32P-radio-labeled probe was prepared and hybridized to different liver total RNA samples from 2-mo-old mice of all different genotypes (Fig. S2 A). The FN RNA from the EDAwt/wt mice (Fig. S2, B and C, lanes 1 and 2) showed 98–99% of the 187 bases protected fragment, indicating almost complete EDA exon exclusion, in complete agreement with previous studies (Tamkun and Hynes, 1983; Kornblihtt et al., 1984). The FN mRNA prepared from EDA+/+ mice showed only the 350-bases protected fragment, indicating constitutive inclusion of the EDA exon (Fig. S2, B and C, lanes 3 and 4) and the absence of alternative splicing of the EDA exon. The FN mRNA from EDA−/− mice showed the complete exclusion of the EDA exon (Fig. S2, B and C, lanes 5 and 6), as expected because the EDA exon was deleted from the genome. Heterozygous mice bearing one EDA+ allele showed values of EDA exon inclusion close to the expected 50% (Fig. S2, B and C, lanes 7, 8, 11, and 12). Analysis in the brain and spleen produced similar
results to those observed in the liver (unpublished data), except for the tissue-specific variation in the basal ratio of EDA inclusion of the EDA<sup>wt</sup> allele.

**Analysis of allele frequencies, offspring, and embryos**

*Mendelian frequency.* Heterozygous crosses (EDA<sup>+/wt</sup> × EDA<sup>+/wt</sup> and EDA<sup>−/wt</sup> × EDA<sup>−/wt</sup>) produced progeny within the expected Mendelian frequency, indicating that there was no intrauterine mortality. We observed 24.9% EDA<sup>−/wt</sup> offspring born from heterozygous EDA<sup>+/wt</sup> mating pairs; and 27.1% EDA<sup>−/wt</sup>, 49.4% EDA<sup>−/wt</sup>, and 23.5% EDA<sup>−/−</sup> from 332 offspring born from heterozygous EDA<sup>−/wt</sup> mating pairs (not significantly different from the expected Mendelian frequency by the χ<sup>2</sup> test; c<sup>2</sup> = 1.355 and c<sup>2</sup> = 0.916, for EDA<sup>−/wt</sup> and EDA<sup>−/−</sup> matings, respectively).

**Litter size and survival rate until weaning.** The average litter size (mean ± SD) was 8.2 ± 2.3 and 7.5 ± 2.2 for each mating group (EDA<sup>+/wt</sup> × EDA<sup>+/wt</sup> and EDA<sup>−/wt</sup> × EDA<sup>−/wt</sup>), respectively. Survival rate after birth, measured until weaning of the litter, was also indistinguishable between the EDA<sup>+/+</sup> and EDA<sup>−/−</sup> mice when compared with that of the EDA<sup>wt/wt</sup> littermates. The average litter size in crosses between homozygous mice of each genotype (EDA<sup>−/−</sup> × EDA<sup>−/−</sup> and EDA<sup>+/+</sup> × EDA<sup>−/−</sup>) was normal, which did not differ significantly from that observed among the EDA<sup>wt/wt</sup> littermates and was within the range observed previously for heterozygous matings. The EDA<sup>+/+</sup> and EDA<sup>−/−</sup> matings produced on average six to seven litters, each spaced by 21–22 d, which is consistent with the normal mouse estrous cycle.

**Somite number.** Embryos (9.5 d.p.c.) derived from timed matings of EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> mice were analyzed for the number of somites and shape. No differences were observed neither in the shape of the embryos nor in the number of somites (number of somites, mean ± SD: EDA<sup>+/+</sup>: 22.7 ± 2.1, n = 10; EDA<sup>−/−</sup>: 21.9 ± 2.5, n = 25; and EDA<sup>wt/wt</sup>: 23.7 ± 2.6, n = 30).

**Gain weight.** Weight gain of 3-mo-old EDA<sup>+/+</sup> (mean ± SD, 34.3 ± 4.7, n = 19) and EDA<sup>−/−</sup> mice (35.0 ± 4.2, n = 25) did not differ significantly from that of EDA<sup>wt/wt</sup> mice weight (33.8 ± 3.4, n = 18).

**Northern blot analysis**

To determine whether the observed decrease in FN protein levels in the EDA<sup>+/+</sup> mice was correlated to a decrease in the mRNA levels, we performed Northern blot analysis with total RNA isolated from the tissues having the higher decrease in FN levels (brain and heart) from 3-mo-old male mice. The 8.0-kb FN messenger was detected by a 711-bp cDNA probe, and the results from the EDA<sup>wt/wt</sup>, EDA<sup>+/+</sup>, and EDA<sup>−/−</sup> mice are shown in Fig. S3 A. No significant variations were observed in RNA samples from the brain and heart of EDA<sup>+/+</sup> mice. To correct for differences in RNA loading, the FN levels were normalized by two housekeeping genes (GAPDH and β-actin) showing similar results (Fig. S3, B–D).

**TUNEL and caspase-3 analysis of skin wounds**

Cell death and apoptosis-specific analysis were performed by the TUNEL and caspase-3 of the 7-d postwounding sections analyzed in Fig. 6 (8 and 16 wounds per genotype for TUNEL and caspase-3 experiments, respectively). We observed no significant differences in the number of positive cells between the different genotypes (Fig. S4).
Supplemental material and methods

Genotyping of progeny by Southern blot and PCR
For PCR analysis, 250–500 ng of genomic DNA was used for 30 cycles of PCR with the FN Nde Fwd and FN Nde Rev primers (5'-CTTCAGGGTGTCTCATAC-3' and 5'-ACCCAGGTGTCTCACTTAG-3', respectively). PCR products were separated in a 1% agarose gel and visualized by ethidium bromide staining (Fig. 2 C).

RNase protection assay
RNase protection analysis was performed as described previously (Ausubel et al., 1998). In brief, the sequence-specific hybridization probe (EDA) was cloned in pBSKII and the antisense radiolabeled RNA was transcribed and gel purified. The RNA samples were hybridized overnight at 45°C with the radioactive labeled antisense transcript, and hybrids were digested with RNases A and T1. Protected fragments were separated on a sequencing 5% acrylamide/urea gels and analyzed by autoradiography. 20 μg of transfer RNA was used as a negative control. All experiments were repeated twice and in duplicates.

Synchronized matings and embryo analysis
Matings were done between homozygous mice of each genotype. One male was left with two females for half an hour in each case for each genotype and sexual intercourse was scored by the presence of a vaginal plug. Gross evaluation of the embryos was done exactly at day 9.5 after copulation. The embryos were carefully dissected under a stereomicroscope, fixed, and the number of somites was counted. Embryos were photographed at the same magnification.

Genotyping of progeny by Southern blot and PCR
Genomic DNA was prepared from tail samples as described previously (Laird et al., 1991). For Southern blot analysis, tail DNA was digested with HindIII and separated electrophoretically on an 0.8% agarose gel, blotted onto a nylon membrane (Z-Probe), and hybridized with a 300-bp radiolabeled DNA fragment containing the EDA exon (Fig. 2 B).

mRNA analysis by radioactive RT-PCR
FN cDNAs were generated using the cDNA synthesis kit (Amersham Biosciences) and alternative splicing of FN mRNA was quantitated via RT-PCR using primers in the exons flanking the EDA exon (FN2 Dir, 5'-ACCATCAC-CCTGTATGCTGTCT-3' and EDA Rev, 5'-CTATGAGTCCTGACACAATC-3') using a 32P-dCTP in the PCR mix. RT-PCR products were analyzed by loading on 5% polyacrylamide gels, and the quantitative analysis of the PCR products was performed using a Cyclone PhosphorImager (Storage Phosphor System; Canberra Packard SRL). The values were corrected for background and G/C contents.

TUNEL analysis of skin wounds
Skin wounds at day 7 after wounding were analyzed with the TUNEL assay kit as described by the manufacturers (In situ Cell Death Detection Kit; Roche). In brief, skin sections were treated with terminal deoxynucleotidyl transferase and incubated with a solution containing the labeled nucleotide that was incorporated into cells having DNA breaks. The images were observed using a confocal microscope (Carl Zeiss MicroImaging, Inc.) and photographed under the same conditions of exposure and magnification (400×).
Caspase-3 analysis of skin wounds

Serial sections of the same wounds used for the TUNEL assay were treated with the anti–caspase-3 antibody (Asp175; Cell Signaling). The samples were incubated with a peroxidase-conjugated secondary antibody by using the EnVision kit (DakoCytomation), and stained with the liquid DAB substrate-chromagen solution (DakoCytomation) as suggested by the manufacturers. The images were observed and photographed at the same magnification (100×).

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