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Inflammatory myofibroblastic tumors (IMTs) are characterized by myofibroblast proliferation and an inflammatory cell infiltrate. Little is known about the molecular pathways that precipitate IMT formation. Here, we report the identification of somatic mutations in *UPF1*, a gene that encodes an essential component of the nonsense-mediated RNA decay (NMD) pathway, in 13 of 15 pulmonary IMT samples. The majority of mutations occurred in a specific region of *UPF1* and triggered *UPF1* alternative splicing. Several mRNA targets of the NMD pathway were upregulated in IMT samples, indicating that the *UPF1* mutations led to reduced NMD magnitude. These upregulated NMD targets included *NIK* mRNA, which encodes a potent activator of NF- κ B. In human lung cells, *UPF1* depletion increased expression of chemokine-encoding genes in a *NIK*-dependent manner. Elevated chemokines and IgE class switching events were observed in IMT samples, consistent with *NIK* upregulation in these tumors. Together, these results support a model in which *UPF1* mutations downregulate NMD, leading to *NIK*-dependent NF- κ B induction, which contributes to the immune infiltration that is characteristic of IMTs. The molecular link between the NMD pathway and IMTs has implications for the diagnosis and treatment of these tumors.

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The nonsense-mediated RNA decay pathway is disrupted in inflammatory myofibroblastic tumors

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Inflammatory myofibroblastic tumors (IMTs) are characterized by myofibroblast proliferation and an inflammatory cell infiltrate. Little is known about the molecular pathways that precipitate IMT formation. Here, we report the identification of somatic mutations in *UPF1*, a gene that encodes an essential component of the nonsense-mediated RNA decay (NMD) pathway, in 13 of 15 pulmonary IMT samples. The majority of mutations occurred in a specific region of *UPF1* and triggered *UPF1* alternative splicing. Several mRNA targets of the NMD pathway were upregulated in IMT samples, indicating that the *UPF1* mutations led to reduced NMD magnitude. These upregulated NMD targets included *NIK* mRNA, which encodes a potent activator of NF- κ B. In human lung cells, *UPF1* depletion increased expression of chemokine-encoding genes in a NIK-dependent manner. Elevated chemokines and IgE class switching events were observed in IMT samples, consistent with NIK upregulation in these tumors. Together, these results support a model in which *UPF1* mutations downregulate NMD, leading to NIK-dependent NF- κ B induction, which contributes to the immune infiltration that is characteristic of IMTs. The molecular link between the NMD pathway and IMTs has implications for the diagnosis and treatment of these tumors.

Introduction

Inflammatory myofibroblastic tumors (IMTs) encompass a wide spectrum of irregular myofibroblastic proliferative growths accompanied by infiltrating immune cells that occur in almost all major organs of the human body (1). While IMTs are rare, constituting only 0.04% of general thoracic surgeries, they often occur in children and can grow to a large size and are often locally invasive (2, 3).

At present, the causes of IMTs are unknown. In this communication, we provide evidence that disruption of an RNA degradation pathway has a causal role in the majority of cases of IMT. This pathway — nonsense-mediated RNA decay (NMD) — degrades subsets of mRNAs and is responsible for regulating developmental events and other biological processes (4). We recently showed that the gene encoding an ATP-dependent RNA helicase essential for NMD, Up-frameshift 1 (*UPF1*; also called *RENT1*), is commonly mutated in pancreatic adenocarcinomas (5), suggesting that mutations in *UPF1* have a causal role in these malignant tumors. In the present study, we report that 86% of IMTs that we examined have *UPF1* mutations. We provide evidence that NMD is strongly perturbed in these tumors, leading to aberrant activation of the NF- κ B pathway and immune responses characteristic of these tumors. These findings afford opportunities to better diagnose and treat these tumors.

Results and Discussion

Somatic *UPF1* mutations in IMTs elicit exon skipping. Based on our previous identification of frequent *UPF1* mutations in pancreatic tumors (5), we screened IMT lung samples to determine whether they also had *UPF1* mutations. Sequence analysis of *UPF1* revealed mutations in IMTs from 13 of 15 patients (Supplemental Table 1 and Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI86508DS1). In total, 41 mutations were detected. All but one of these 41 mutations were clustered at 11 sites in exon 10 and intron 10 (Supplemental Figure 1B and Supplemental Table 1). Mutations at 8 of these 11 sites were found in more than one tumor (Supplemental Figure 1B). The mutations in the 13 IMTs were not detected in disease-free lung tissues and thus were somatic in origin (data not shown).

UPF1 exon 10 and intron 10 are unusually short (169 and 85 nt, respectively) and have a high GC content (59% and 73%, respectively), both of which are features that can weaken RNA splicing (6). This raised the possibility that intron 10 splicing depends on local splicing enhancers (7) and that these elements are disrupted by the mutations we identified in the *UPF1* gene, leading to aberrant *UPF1* splicing. To test this possibility, we constructed a *UPF1* minigene containing the relevant region of *UPF1* (Figure 1A). When transfected into HEK293 cells, this minigene was normally spliced, as shown by direct sequencing of the only band observed by RT-PCR analysis (Figure 1B). We then introduced the mutations found in IMTs (Figure 1A and Supplemental Figure 2A). This caused reduced expression of the normally spliced mRNA and the appearance of an alternatively spliced transcript lacking exons 10 and 11 (Figure 1B and Supplemental Figure 2B).

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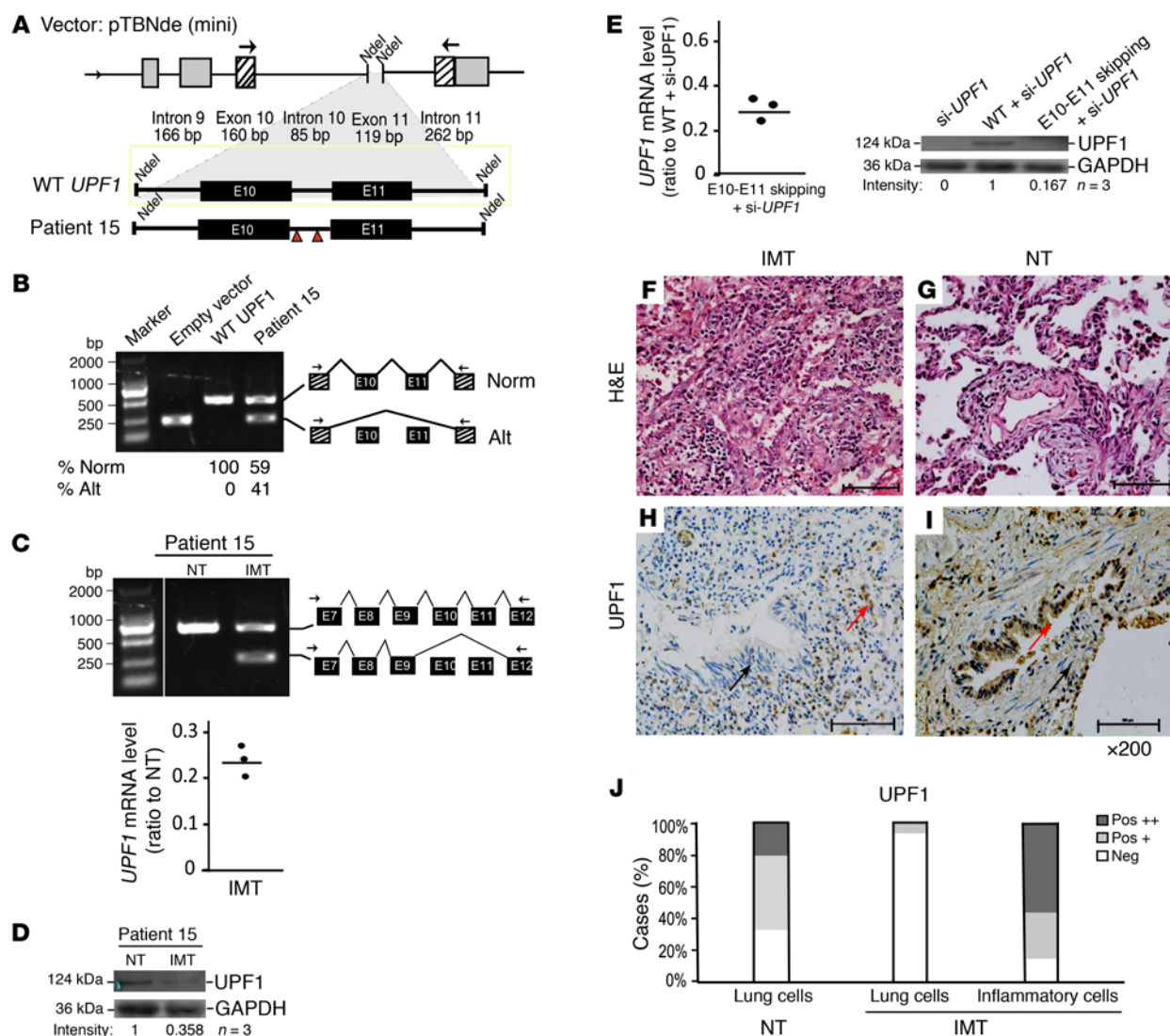


Figure 1. IMT-specific *UPF1* mutations disrupt splicing of *UPF1* mRNA and reduce *UPF1* protein levels. (A) Patient 15 mutations were inserted into the wild-type human *UPF1* minigene as previously described (5). (B) RT-PCR analysis of HEK293 cells transfected with the constructs shown in A (primer locations are indicated by the arrows) ($n = 5$). Direct sequencing of the large (518 bp) and small (239 bp) bands indicated that they correspond to normally spliced (Norm) and exon-skipped transcripts (Alt), respectively. The numbers below the gel are the average values from 5 independent transfections. (C) RT-PCR (upper) and RT-qPCR (lower) analysis of endogenous *UPF1* expression in IMT lung tissue and normal lung tissue (NT) from patient 15 (direct sequencing results correspond to the indicated schematics [$n = 5$]). (D) Western analysis of patient 15 ($n = 3$). Because of the large size difference, *UPF1* and the loading control, GAPDH, were assayed on different percentage polyacrylamide gels (but with the same amount of sample loaded); the lanes shown were run on the same gel but were noncontiguous. (E) RT-qPCR (left) and Western (right) analysis of HEK cells transfected with the indicated *UPF1* expression vectors ($n = 3$). GAPDH is the loading control ($n = 3$). Western analysis performed in parallel gels was done as in D. (F and G) H&E staining of patient 15 tissue. (H and I) IHC staining of patient 15 tissue ($\times 200$). Red arrows indicate strong staining; black arrows indicate weak or negative staining ($n = 6$). (J) Quantification of H and I. Pos +, positive staining of greater than 20% of cells; Pos ++, positive staining of greater than 50% of cells.

To examine endogenous *UPF1* splicing, we analyzed the IMT tumor from patient 15, which was frozen, providing sufficiently high quality RNA for analysis. RT-PCR analysis revealed an alternatively spliced *UPF1* transcript identical to that generated by the mutant minigene in IMT tissue (Figure 1C). This alternatively spliced *UPF1* transcript was not present in normal lung tissue (NT) from the same patient. Consistent with the reduced level of *UPF1* mRNA in IMT (Figure 1C), the *UPF1* protein level was also reduced (Figure 1D). To investigate the mechanism, we generated a *UPF1* cDNA construct lacking exons 10 and 11. Transfection of this mutant construct into HEK cells depleted of endogenous *UPF1*

revealed that it was expressed at a significantly lower steady-state RNA level than was the full-length *UPF1* construct (Figure 1E), an effect that was mirrored at the protein level (Figure 1E). The simplest explanation for these findings is that the loss of exons 10 and 11 destabilizes *UPF1* RNA.

As an independent means to analyze *UPF1* protein expression levels, we examined IMT tissue sections, which unlike NT, had not only alveolar epithelial cells but also inflammatory cells (Figure 1, F and G, and Supplemental Figure 3), as is characteristic of IMTs (1, 3). Immunohistochemical (IHC) analysis with a highly specific *UPF1* antibody showed that normal alveolar epithelial cells expressed

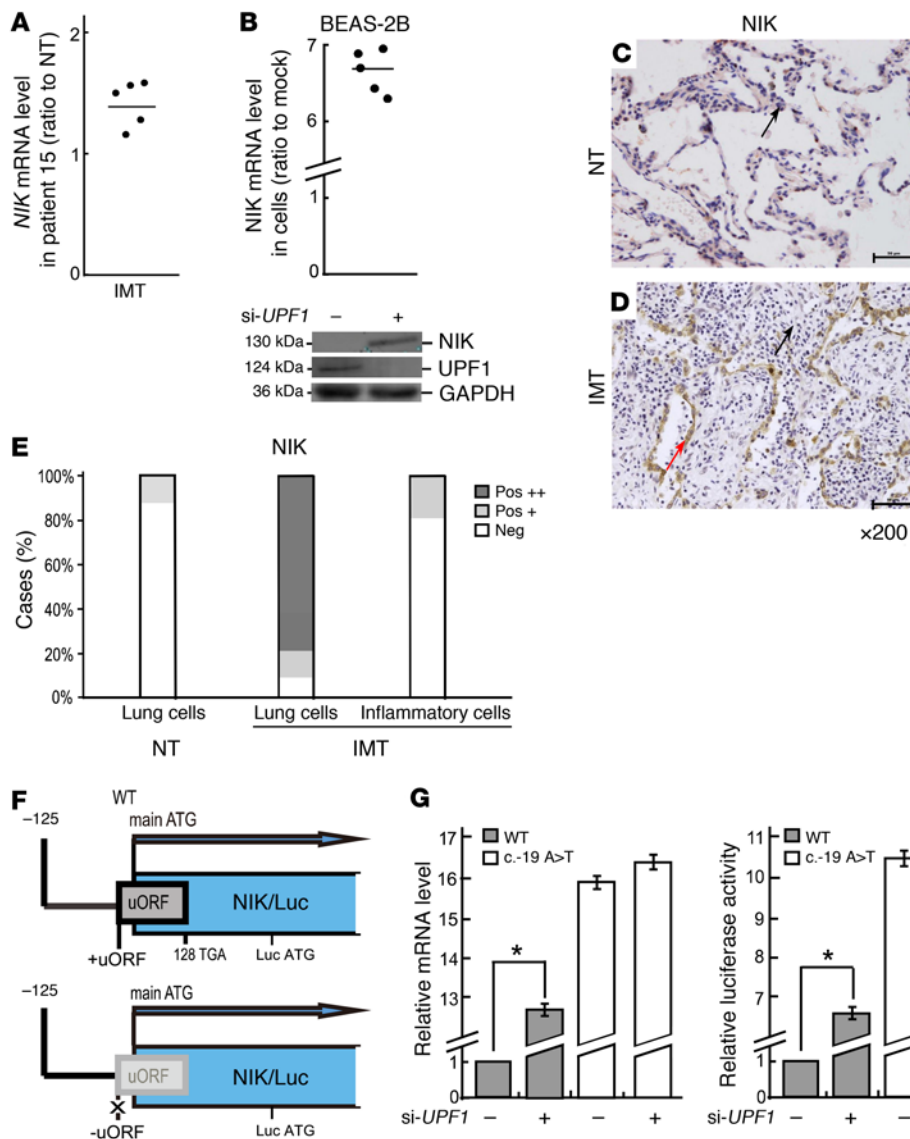


Figure 2. NIK expression is upregulated in IMTs harboring *UPF1* mutations. (A) RT-qPCR analysis of IMT lung tissue and normal lung tissue (NT) from patient 15 ($n = 5$). (B) RT-qPCR (top [$n = 5$]) and Western (bottom [$n = 3$]) analysis of BEAS-2B cells depleted of *UPF1*. GAPDH is the loading control. Western analysis performed in parallel gels was done as in Figure 1D. (C and D) IHC staining in patient 15 tissue ($\times 200$). Red arrow, strong staining; black arrows, weak or negative staining ($n = 6$). (E) Quantification of C and D. Pos +, positive staining in greater than 20% of cells; Pos ++, positive staining in greater than 50% of cells. (F) *NIK* 5' UTR cloned upstream of a luciferase (*Luc*) reporter gene. The 147-nt uORF with or without a mutated start ATG is shown as a gray box. (G) Relative *Luc* mRNA levels (left [$n = 5$]) and *Luc* enzymatic activity (right [$n = 5$]) from transfected BEAS-2B cells. For G, the values are the average fold change (mean \pm SEM) from 5 independent experiments (Student's *t* test, $*P < 0.05$). 18S mRNA was used as the endogenous control.

high levels of *UPF1*, while most patient IMT alveolar epithelial cells exhibited little or no *UPF1* expression (Figure 1, H–J, Supplemental Figure 3, and Supplemental Table 2). The only exceptions were the epithelial cells in the two IMTs lacking *UPF1* mutations and the IMT with a missense *UPF1* mutation (in exon 13), all of which expressed high levels of *UPF1* (Supplemental Figure 3). In contrast with the alveolar epithelial cells, the inflammatory cells in IMTs had high *UPF1* antibody reactivity regardless of *UPF1* mutation status (Supplemental Figure 3 and Supplemental Table 2). These findings suggest that *UPF1* mutations occur specifically in the alveolar epithelial cells in IMTs, leading to reduced *UPF1* protein levels in these cells.

We next examined the functional consequences of the single *UPF1* mutation we identified outside of the exon 10–11 region (c.1796A>T p.K599M from patient 13; Supplemental Table 1). The strong *UPF1* IHC staining in this patient sample (Supplemental Table 2) suggested that this mutation does not destabilize *UPF1* RNA, and raised the alternative possibility that it disrupts *UPF1* activity. In support, we found that exogenous *UPF1* harboring the missense mutation corresponding to patient 13 failed to rescue NMD activity in *UPF1*-depleted cells (as it did not significantly

decrease the level of an NMD-sensitive NS39 β -globin mRNA), indicating that this mutant lacked detectable NMD function. Furthermore, this mutant conferred dominant-negative activity, as it reduced the activity of exogenous WT *UPF1* (Supplemental Figure 4). Together with our analysis of the other *UPF1* mutants described above, this suggested that two types of mutations are employed to reduce *UPF1* activity in IMTs: those that generate a dominant-negative *UPF1* or an unstable form of *UPF1* mRNA.

IMTs have disrupted NMD and elevated expression of the pro-inflammatory molecule *NIK*. The dramatically reduced level of *UPF1* expression in most IMTs raised the possibility that NMD is depressed in IMTs, leading to upregulation of NMD target transcripts. In agreement with this hypothesis, we found that 11 endogenous NMD substrates were upregulated in the patient 15 IMT sample relative to NT from the same patient (Supplemental Table 3). We next considered that perturbed NMD would allow mRNAs encoding proinflammatory proteins to be stabilized, leading to the immune infiltration characteristic of IMTs. We identified mitogen-activated protein kinase kinase kinase 14 (MAP3K14 or *NIK*) as a good candidate, as it is an activator of the proinflammatory NF- κ B

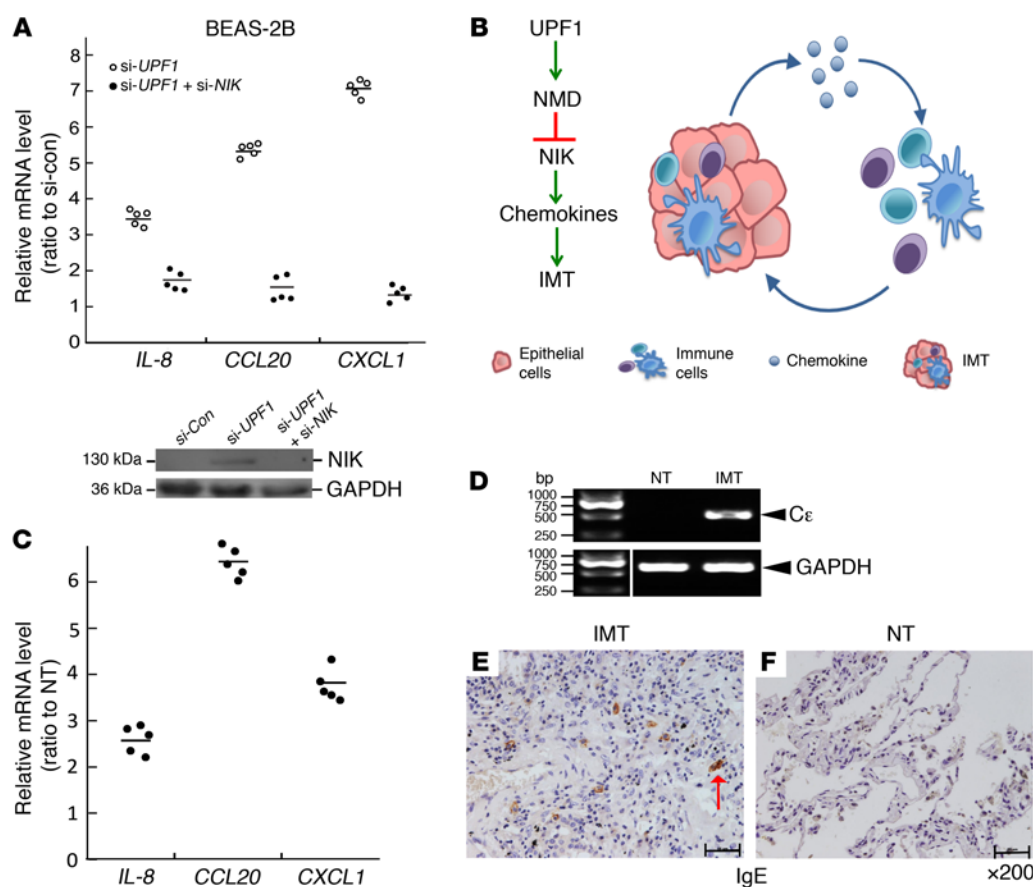


Figure 3. Evidence for a UPF1-based circuit that elevates chemokine and IgE expression.

(A) RT-qPCR (top [$n = 5$]) and Western (bottom [$n = 3$]) analysis of BEAS-2B cells depleted of *UPF1* and/or *NIK*. *18S* mRNA and GAPDH are the endogenous controls. (B) Model depicting how NMD deficiency causes IMT formation. (C) RT-qPCR analysis of IMT lung tissue and normal lung tissue (NT) from patient 15 ($n = 5$). (D) RT-PCR analysis of patient 15 tissue ($n = 5$). Direct sequencing of the band indicated that it is a *Cε* germline transcript (exons 1 and 3; 487 bp). GAPDH is the loading control. The lanes were run on the same gel but were noncontiguous. (E and F) IHC analysis of patient 15 ($\times 200$). The red arrow, IgE-positive cell ($n = 3$).

signaling pathway (8). In support of its potential role, we found that *NIK* mRNA was upregulated in the patient 15 IMT relative to NT (Figure 2A) and in response to depletion of *UPF1* in the normal lung epithelial cell line BEAS-2B (Figure 2B, upper). Upregulation of *NIK* in BEAS-2B cells was confirmed at the protein level (Figure 2B, lower) and in IMT tissue harboring *UPF1* mutations, as detected by IHC analysis (Figure 2, C–E, and Supplemental Figure 5). In contrast, neither of the two IMT tumors lacking detectable *UPF1* mutations had upregulated *NIK* protein expression (Supplemental Table 2). Upregulated *NIK* expression was specific to lung tumor cells, as the local infiltrating leukocytes had low/no *NIK* expression in all patients (Supplemental Figure 5 and Supplemental Table 2).

NIK mRNA was previously suggested to be an NMD substrate (9, 10), and so we assessed the mechanism by which *NIK* mRNA is degraded by NMD. Stop codons terminating short ORFs upstream of the protein-coding ORF can elicit NMD (4). Thus, we considered the possibility that the upstream ORF (uORF) present in the 5' UTR of *NIK* mRNA is responsible for it being degraded by NMD. In support of this, insertion of the *NIK* 5' UTR in front of a luciferase (Luc) reporter (Figure 2F) led to increased reporter expression in *UPF1*-depleted cells, as measured at the RNA level or by Luc activity (Figure 2G). Mutation of the uORF start ATG by replacing the "A" with a "T" (–19A>T) (Figure 2F) completely abolished this reporter upregulation (Figure 2G). We conclude that *NIK* mRNA is an NMD substrate by virtue of its possessing a uORF.

Stabilized *NIK* elevated chemokine gene expression and IgE production. *NIK* is a serine/threonine protein kinase that stimulates NF- κ B activity (8). Given that a hallmark of NF- κ B activation is the

generation of chemokines (11), this raised the possibility that the upregulation of *NIK* in IMTs has a causal role in these inflammatory tumors. In support of this possibility, we found that *UPF1* depletion dramatically increased *IL-8*, *CCL20*, and *CXCL1* mRNA expression in the BEAS-2B lung cell line (Figure 3A). This upregulated chemokine expression resulted, at least in part, from upregulated *NIK* expression, as it was largely abolished when we prevented *NIK* upregulation using RNAi (Figure 3A). Together, these data support a model in which NMD normally serves to suppress the chemokine response by destabilizing *NIK* mRNA (Figure 3B). When NMD is perturbed, this suppression is lifted, leading to high *NIK* expression and the generation of high levels of chemokines, thereby triggering immune cell infiltration, the hallmark of IMTs (1). As evidence that this model applies in vivo, RT-qPCR analysis revealed upregulated expression of the chemokine genes *IL-8*, *CCL20*, and *CXCL1* in the patient 15 IMT relative to NT (Figure 3C).

Another property of *NIK* is its ability to promote B cell proliferation and induce B cells to differentiate into IgE-secreting plasma cells (12, 13). The commitment of a B cell to isotype class switch to an IgE-producing cell requires induction of *Cε* germline transcription (14, 15). We thus assessed whether IMTs have upregulated germline *Cε* transcripts. Using an RT-PCR assay, we observed a 0.5-kb band indicative of *Cε* transcription, which we confirmed was a germline *Cε* transcript by direct sequencing (Figure 3D). Consistent with activation of *Cε* transcription and rearrangement, IHC analysis showed that patient 15 had many IgE-positive cells (Figure 3E), whereas normal lung tissue from the same patient did not (Figure 3F). The IgE-positive cells in the IMT tissue had the morphology of plasma or mast cells.

Our data support a model in which NMD normally serves to confer tight regulation on NIK abundance in order to place cells below the threshold required for NF- κ B pathway induction and inappropriate immune activation. By analogy, NMD has been shown to raise the threshold for triggering the unfolded protein response, thereby preventing programmed cell death in response to innocuous stimuli (16, 17). Interestingly, the vast majority of the mutations that we observed in IMTs were clustered in one of the same two regions of the *UPF1* gene as in adenocarcinoma pancreatic cancer (the exon 10-intron 10-exon 11 region; Supplemental Figure 1B). Indeed, 7 of the pulmonary IMTs had *UPF1* mutations at precisely the same site as in some ASC tumors (IVS10+32C>T and IVS10-25C>T). We speculate that the aberrations in NMD caused by genetic alterations in the *UPF1* gene lead to different disease outcomes, depending on the cell type. In premalignant glandular pancreatic cells, this may lead to reprogramming towards a squamous, more malignant state (5). In lung epithelial cells, NMD perturbation may drive a proinflammatory response, as suggested by our results herein.

Our finding that *UPF1* mutations are a common molecular alteration associated with IMTs suggest that *UPF1* mutations and NMD activity may be useful diagnostic markers for this disease. New diagnostic markers are critical, as IMTs are often misdiagnosed as malignant tumors (1, 2). Further, our results described herein suggests that therapies directed at restoring the NMD pathway and NIK expression have the potential to ameliorate or cure the symptoms of IMT and related diseases, including IgG4-related disease (IgG4-RD), an autoimmune disorder that resembles IMTs (1, 3). New therapies are desired, as current methods to treat IMTs and IgG4-RD, including surgery and antiinflammatory agents, are invasive or have undesirable side effects (2, 3).

Methods

Subjects. We evaluated a set of IMTs from 15 patients, provided by the First Affiliated Hospital of Bengbu Medical College. Among them, 14

specimens were formalin-fixed paraffin-embedded (FFPE) tissue sections, and one IMT of patient 15 was frozen immediately after collection. All patients provided written informed consent.

Statistics. The program ESE Finder was used to locate several predicted ESEs and ISEs in the regions of the *UPF1* gene. Pairwise comparisons were performed by 2-tailed Student's *t* test using Excel software (Microsoft). Data are expressed as the mean \pm SEM.

Study approval. All aspects of the study were approved by the Bengbu Medical College and Tongji University School of Medicine.

For DNA/RNA sequencing and analysis, constructs, cell culture and transfections, and protein analysis, see Supplemental Methods.

Author contributions

CL, YW, GY, and TP performed UPF1 screening. JL and AH constructed wild-type and mutant mini-gene expression vectors. JL, FS, SL, and YW performed IHC, Western, and RNA analyses. FS, CS, YJ, and CY provided pathology samples, clinical data, and expertise on IMTs. JL and TP prepared figures and tables and wrote the manuscript. YL and MFW designed and supervised the project and assisted in writing the manuscript.

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