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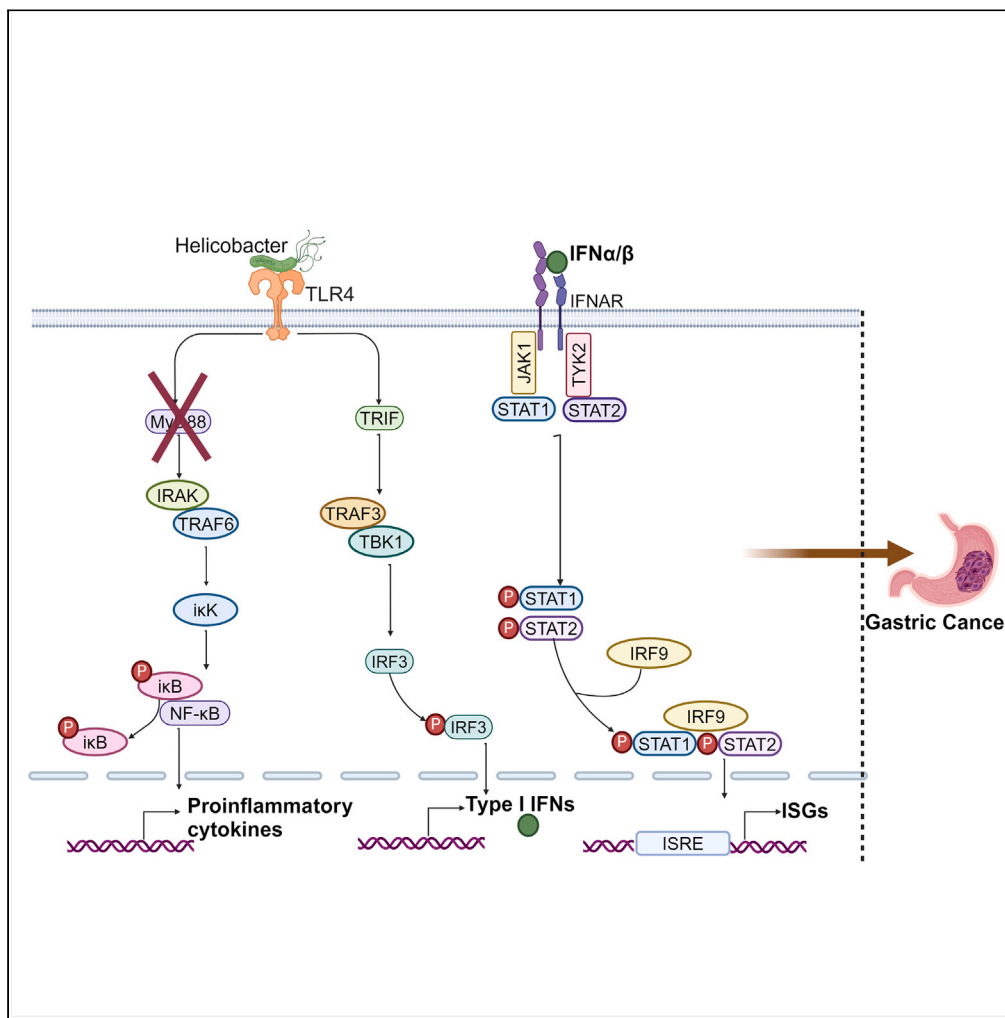
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Article

TRIF-IFN-I pathway in *Helicobacter*-induced gastric cancer in an accelerated murine disease model and patient biopsies



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Highlights

Helicobacter-induced gastric cancer (GC) progresses rapidly in absence of MyD88

Absence of TRIF decreased *Helicobacter* induced disease pathology

ISGs upregulated in *Helicobacter* infected *Myd88*^{-/-} mice and GC patients

TRIF-IFN-I pathway plays pivotal role in development of *Helicobacter*-induced GC



Article

TRIF-IFN-I pathway in *Helicobacter*-induced gastric cancer in an accelerated murine disease model and patient biopsies

Prerna Bali,¹ Ivonne Lozano-Pope,¹ Jonathan Hernandez,¹ Monica V. Estrada,² Maripat Corr,¹ Michael A. Turner,^{3,4} Michael Bouvet,^{2,3,4} Christopher Benner,¹ and Marygorret Obonyo^{1,2,5,*}

SUMMARY

***Helicobacter pylori* (*H. pylori*) infection is a known cause of many digestive diseases, including gastritis, peptic ulcers, and gastric cancer. However, the underlying mechanisms by which *H. pylori* infection triggers these disorders are still not clearly understood. Gastric cancer is a slow progressing disease, which makes it difficult to study. We have developed an accelerated disease progression mouse model, which leverages mice deficient in the myeloid differentiation primary response 88 gene (*Myd88*^{-/-}) infected with *Helicobacter felis* (*H. felis*). Using this model and gastric biopsy samples from patients, we report that activation of the Toll/interleukin-1 receptor (TIR)-domain-containing adaptor inducing interferon- β (TRIF)-type I interferon (IFN-I) signaling pathway promotes *Helicobacter*-induced disease progression toward severe gastric pathology and gastric cancer development. Further, results implicated downstream targets of this pathway in disease pathogenesis. These findings may facilitate stratification of *Helicobacter*-infected patients and thus enable treatment prioritization of patients.**

INTRODUCTION

Helicobacter pylori (*H. pylori*) infection is the main risk factor for development of gastric cancer, thus this bacterium is classified as a group I carcinogen.^{1,2} Gastric cancer, which is often recalcitrant to therapy, remains a significant global health burden and a leading cause of cancer-related deaths worldwide.^{3,4} Half of the world's population is infected with *H. pylori*, contributing significantly to the disease burden.^{2,5} Infection with *H. pylori* causes chronic inflammation, leading to the development of preneoplastic lesions (atrophy, intestinal metaplasia, dysplasia), and ultimately to gastric cancer, a progression known as the Correa's cascade.⁶ We previously demonstrated that mice deficient in myeloid differentiation primary response gene 88 (*MyD88*, *Myd88*^{-/-}) infected with *H. felis* rapidly developed gastric pathology progressing to high-grade gastric dysplasia,⁷ a condition considered as gastric cancer *in situ*.^{8,9} Mechanisms underlying this phenotype were not fully understood, but our data indicated that the loss of *MyD88* led to hyperactivation of the alternate Toll/interleukin-1 receptor (TIR) domain-containing adaptor inducing interferon- β (TRIF)-dependent pathway, accelerating gastric cancer progression in response to *H. felis* infection.^{7,10} A recent study citing our work revealed that the Toll-like receptor 4 (TLR4)-TRIF-type I interferon (IFN-I) pathway, via the production of IFN- γ , was important for *H. suis*-induced disease development in a mucosa-associated lymphoid tissue (MALT) lymphoma mouse model.¹¹ *MyD88* and TRIF are the main adaptor proteins that mediate TLRs signals.¹² TRIF signaling branches into two downstream pathways, one that shares effectors with *MyD88* and another that is *MyD88*-independent, (TRIF/IRF3, IFN regulatory factor 3),^{13–15} leading to the induction of IFN-I genes and IFN-inducible genes.

In this study, we utilized our accelerated animal model for gastric cancer and clinical gastric specimens from patients to establish that the activation of the TRIF-IFN-I signaling pathway promotes *Helicobacter*-induced disease progression, leading to severe gastric pathology and ultimately resulting in the development of gastric cancer. We made the significant observation that these changes in the gastric epithelium begin early, just one month after *Helicobacter* infection, and become even more pronounced at 3 months post-infection. This pathway may therefore be an attractive target to intercept the neoplastic process and prevent the development of gastric cancer.

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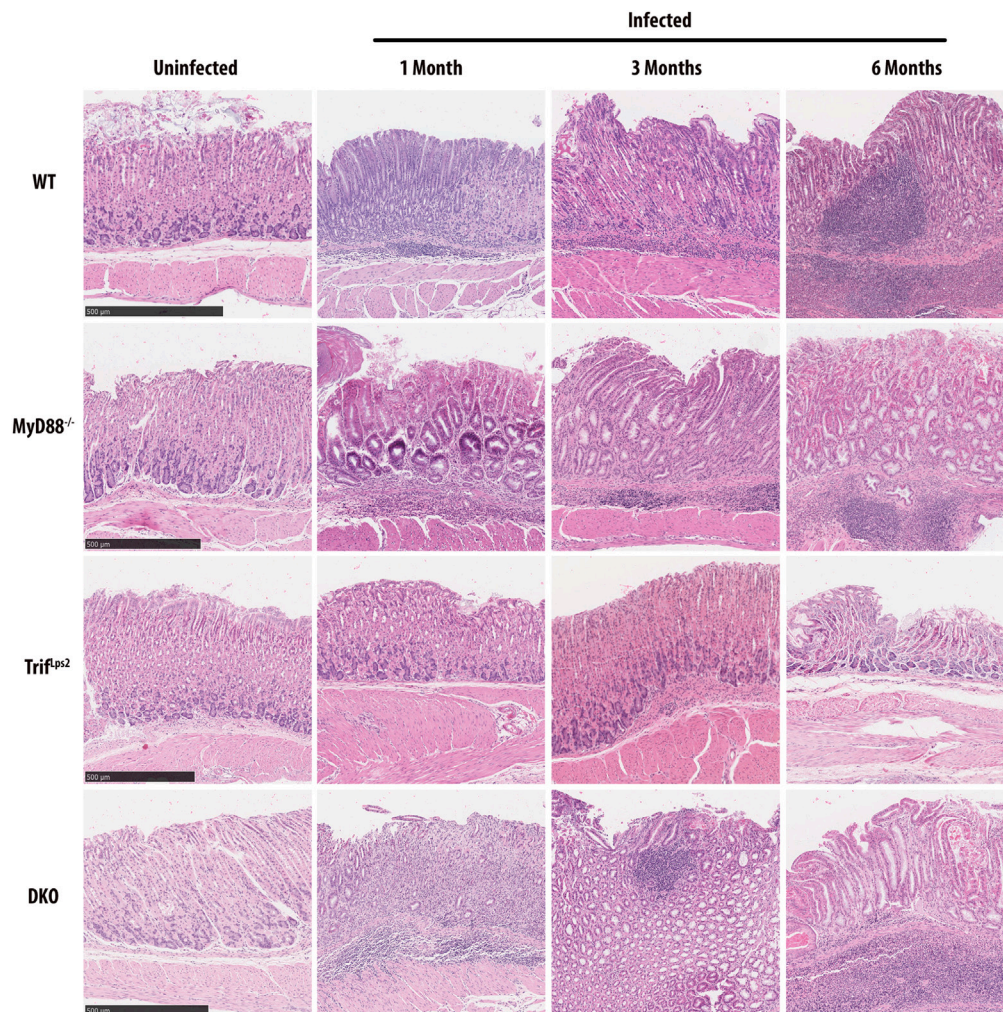


Figure 1. Representative images of hematoxylin & eosin-stained mouse stomach tissue sections

WT, *Myd88*^{-/-}, *Trif*^{Δps2}, and double deficient (*Myd88*^{-/-} and *Trif*^{Δps2}, DKO) mice were infected with *H. felis* for 1 month (WT n = 8; *Myd88*^{-/-} n = 8; *Trif*^{Δps2} n = 8; DKO n = 8), 3 months (WT n = 8; *Myd88*^{-/-} n = 8; *Trif*^{Δps2} n = 8; DKO n = 7), or 6 months (WT n = 7; *Myd88*^{-/-} n = 8; *Trif*^{Δps2} n = 8; DKO n = 8). Scale bar, 500 μm.

RESULTS

Preneoplastic gastric lesions in response to *Helicobacter* infection in mice

Our previous *in vivo* data using *Myd88*^{-/-} mice demonstrated that infection with *H. felis* for 25 or 47 weeks resulted in rapid disease progression with severe gastric pathology.⁷ However, it remained unclear when gastric neoplastic changes were initiated following *H. felis* infection. To address this, we profiled the gastric epithelium at earlier time points post-infection with *H. felis*. Wild type (WT) and *Myd88*^{-/-} mice in the C57BL/6 background were infected with *H. felis* for 1, 3, or 6 months. Considering the role of the TRIF signaling pathway in *H. felis*-induced disease progression from our previous study,⁷ we also infected *Trif*^{Δps2} and double deficient (*Myd88*^{-/-} and *Trif*^{Δps2}, DKO) mice with *H. felis* for 1, 3, and 6 months. All infected mice were colonized, albeit with significant differences in colonization level among strains (Figure S1). Gastric histopathological changes were minimal in *Trif*^{Δps2} mice in response to infection with *H. felis* compared to WT, *Myd88*^{-/-} and DKO mice (Figure 1). In contrast, gastric precancerous lesions developed and progressed more rapidly, with significantly increased severity in *H. felis*-infected *Myd88*^{-/-} mice compared to *H. felis*-infected WT, *Trif*^{Δps2}, and DKO mice. To closely define gastric epithelial changes during *H. felis* infection, we scored histopathology parameters, including inflammation, epithelial defects, oxyntic gland atrophy, hyperplasia, intestinal metaplasia, and dysplasia, on H&E-stained stomach tissue samples by a pathologist blinded to the study. Differences in histopathologic scores of these parameters were evident in *Myd88*^{-/-} mice at 1 month following *H. felis* infection, compared to WT, *Trif*^{Δps2}, and DKO mice, with disease severity increasing with time (Figures S2A–S2F). There were no significant gastric histopathologic changes in all uninfected mice (Figures S2G–S2L). Notably, the earliest significant gastric epithelial changes in all infected mice occurred at 3 months. Consequently, we performed comparative analyses in mice at 3 months following *H. felis* infection. Information gathered at this early phase of infection is ideal

for identifying early targets for disease interception as the disease progresses. This is of central importance due to the lack of characteristic symptoms during early disease stages, often leading to diagnosis at a terminal stage with unfavorable outcomes for patients.

Helicobacter induced high expression of interferon-stimulated genes (ISGs) in the absence of MyD88 signaling in mice

The histopathology data revealed significant changes in the gastric epithelium of *Myd88*^{-/-} mice, including preneoplastic lesions, following *H. felis* infection as early as 3 months, compared to WT, *Trif*^{ps2}, and DKO mice. To identify possible molecular mechanisms driving this disease progression in the absence of MyD88 signaling, we performed RNA-seq to compare gene expression in mouse stomachs between WT and *Myd88*^{-/-} mice after three months of *H. felis* infection. We used mouse stomach tissues from 19 mice for RNA-seq, 8 uninfected (WT n = 2; *Myd88*^{-/-} n = 2; *Trif*^{ps2} n = 2; DKO n = 2), and 11 infected (WT n = 3; *Myd88*^{-/-} n = 2; *Trif*^{ps2} n = 3; DKO n = 3) mice. This analysis showed that many of the significant differentially expressed genes between *Myd88*^{-/-} and WT mice were ISGs, which were highly expressed in gastric tissue samples from *Myd88*^{-/-} mice (log₂ fold change >1, adj. p value <0.05, n = 221, Figure 2A) that exhibited severe gastric pathology. These ISGs were not highly expressed in *Trif*^{ps2} (Figure S3A) or DKO (Figure S3B) mice. Pathway enrichment analysis revealed that these ISGs were the dominant upregulated genes in *Myd88*^{-/-} mice in response *H. felis* (Figure 2B), and their expression was closely correlated with severe gastric pathology, suggesting their involvement in disease pathogenesis. In contrast, the interferon pathway was not enriched in both stomach tissue samples from *Trif*^{ps2} (Figure S4A) or DKO (Figure S4B) mice. Notably, specific ISGs that were upregulated in *Myd88*^{-/-} mice in response *H. felis* infection included indoleamine 2,3-dioxygenase 1 (*Ido1*), guanylate binding protein 2 (*Gbp2*), transcription factor interferon regulatory factor 1 (*Irf1*), beta-2 microglobulin (*B2m*), Z-DNA binding protein 1 (*Zbp1*), and chemokine (C-X-C motif) ligand 11 (*Cxcl11*), *Cxcl9*, *Ccl8*, MX dynamin-like GTPase 1 (*Mx1*), interferon gamma induced GTPase (*Igtp*), proteasome subunit beta 8 (*Psmβ8*), and immunity-related GTPase family M member 1 (*Irgm1*) (Figure 3A). These ISGs were specifically induced in the gastric tissue from *H. felis*-infected *Myd88*^{-/-} mice and were not observed in gastric tissue samples from *H. felis*-infected WT, *H. felis*-infected *Trif*^{ps2}, *H. felis*-infected DKO, or uninfected mice. We further observed that the upregulation of these ISGs in response to *H. felis* infection was highly specific. ISGs are typically induced through activation of the IFN-I pathway.¹⁶ Comparing upregulation of IFN-I targets with that of nuclear factor-kappa B (NF-κB) targets in response to *H. felis* infection, we found minimal differences in the upregulation of NF-κB targets between mice, as exemplified by the NF-κB target *Nfκbia*. However, the upregulation of IFN-I targets (i.e., ISGs) were exclusively observed in the absence of MyD88 signaling (*Myd88*^{-/-} mice) (Figure 3B). This specific association of MyD88-deficiency with a heightened ISG profile strongly suggests the activation of the TRIF-IFN-I signaling pathway.

To understand the possible pathogenic mechanism of these ISGs in *Helicobacter*-induced disease, we performed promoter motif enrichment analysis of genes induced in the stomachs of WT, *Myd88*^{-/-}, *Trif*^{ps2}, and DKO mice following *H. felis* infection. In line with the upregulation of ISGs in our *H. felis*-infected *Myd88*^{-/-} model, this analysis revealed that the IFN stimulated-response element (ISRE) motif was the single most enriched motif in promoters of genes upregulated in *H. felis*-infected *Myd88*^{-/-} mice (Figure 4). In contrast, this motif was not enriched *Trif*^{ps2} and DKO mice. This provides strong evidence that IFN-I regulatory pathways and transcription factors play an important role in the severe disease pathology observed in these mice in response to *H. felis* infection. Since ISGs mainly mediate IFN effects,¹⁶ these findings suggest a significant role of the TRIF-IFN-I pathway in driving severe gastric pathology in response to *H. felis* infection.

Decreased Helicobacter-induced disease pathology in the absence of TRIF signaling in mice

Following infection with *H. felis* for 3 months, histopathology scoring of gastric tissue from WT, *Myd88*^{-/-}, *Trif*^{ps2}, and DKO mice revealed that inflammation was induced in all mouse groups, but it was significantly less severe in *Trif*^{ps2} mice compared to WT, *Myd88*^{-/-}, and DKO mice (Figure 5A). While the disease progressed in WT, *Myd88*^{-/-}, and DKO mice, leading to oxyntic gland atrophy, *Trif*^{ps2} mice did not show evidence of this pathology (Figure 5B). Oxyntic atrophy, representing the loss of parietal cells, is known to be a pivotal step in the progression of lesions, including intestinal metaplasia and dysplasia, eventually leading to gastric cancer.^{17,18} Importantly, in the absence of TRIF signaling, *Helicobacter*-induced inflammation did not progress to precancerous lesions, including intestinal metaplasia (Figure 5C) and dysplasia (Figure 5D). The DKO mice with double deficiency in both TRIF and MyD88 signaling showed an intermediate dysplasia score. These findings indicate that deficiency in TRIF signaling mitigates the severe gastric pathology observed in *Myd88*^{-/-} mice in response to *H. felis* infection, corroborating our previous studies indicating that the TRIF pathway plays a key role in the development of severe disease pathology. Additionally, a recent study citing our previous work suggested that TLR4-TRIF signaling was important in a MALT lymphoma model involving infection of mice with *H. suis*.¹¹ Together, our current study and previous work indicate that the TRIF signaling pathway, and consequently downstream activation of IFN-I pathway, may be responsible for early development and accelerated disease progression. This supports the role of downstream target genes of this pathway in promoting disease progression, as we observed a significant association of high-grade dysplasia with the upregulation of ISGs.

Elevated ISG expression in patient gastric tissue samples is closely associated with progression of gastric cancer

The investigation of ISGs in previous studies of *H. pylori*-induced disease provided valuable insights. Upon reanalyzing microarray data (Gene Expression Omnibus, GEO; Accession number GSE27411),¹⁹ which included gastric biopsies of *H. pylori*-infected and uninfected patients with atrophic gastritis, we observed elevated levels of ISGs ((2'-5'-oligoadenylate synthase 2 (*Oas2*), *B2m*, *Gbp2*, *Mx1*, *Irf1*, *Ido1*, and *Cxcl9*) in gastric tissue samples from patients with *H. pylori*-induced chronic atrophic gastritis (Figure 6) similar to our findings observed in *Myd88*^{-/-} mice following *H. felis* infection. As chronic atrophic gastritis is considered the initial step in the multi-step pre-neoplasia progression cascade,²⁰ the overexpression of ISGs could signify a transition toward severe disease pathology.

To further validate these findings, we examined available gastric samples obtained from the UCSD Biorepository. Our analysis revealed significant upregulation of several ISGs (*Oas2*, *B2m*, ISG product 15 (*Isg15*), *Gbp2*, *Mx1*, *Irf1*, *Ido1*, and *Cxcl9*) in human gastric cancer biopsy

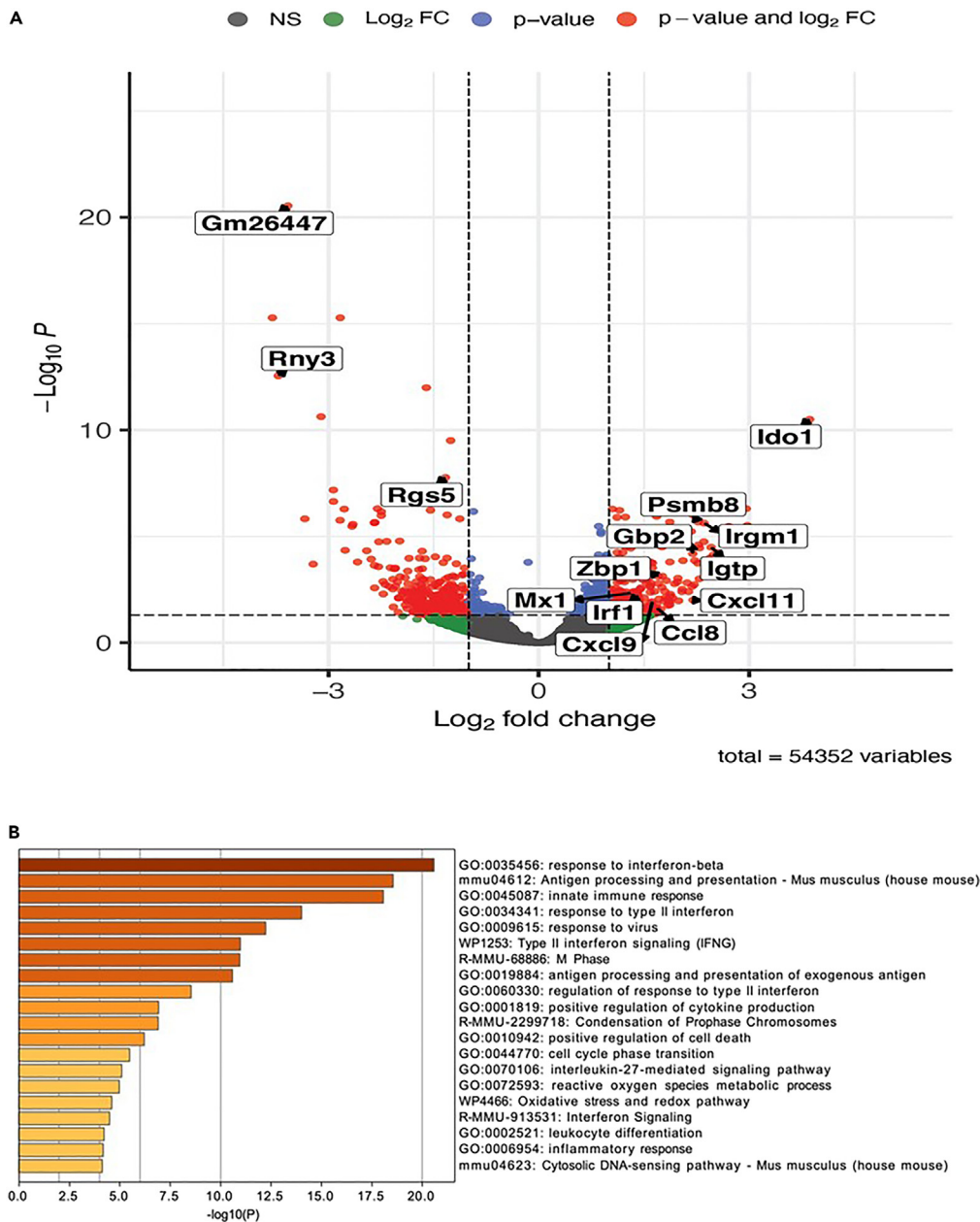


Figure 2. Genes expression and pathway analysis in mouse gastric tissue

(A) Volcano plot shows expression of genes following three months of infection with *H. felis* in WT and *Myd88*^{-/-} mice. The plot depicts the log₂ fold change and adjusted log₁₀ p values for each gene when comparing gene expression values between *felis*-infected WT and *H. felis*-infected *Myd88*^{-/-} mice gastric tissue. (B) Metascape pathway enrichment analysis of the top 20 pathways and their adjusted log₁₀ p value enrichment values in genes that were upregulated in *H. felis*-infected *Myd88*^{-/-} mice gastric tissue samples relative to *H. felis*-infected WT gastric tissue samples.

samples, mirroring the findings observed in gastric tissue samples from *H. felis*-infected *Myd88*^{-/-} mice (Figure 7). This validation of highly expressed ISGs between *H. felis*-infected *Myd88*^{-/-} mice and human gastric tissue samples from *H. pylori*-infected and gastric cancer patients strengthens the evidence for these ISGs being involved in disease progression to gastric cancer.

Validation of mouse data in gastric cancer patient biopsy samples using Kaplan-Meier survival analysis

In our previous study, we found that mice deficient in *Myd88* showed rapid progression of gastric pathology upon *H. felis* infection, with more severe gastric pathology including severe gland atrophy, hyperplasia, intestinal metaplasia, and gastric dysplasia than WT mice.⁷ Two recent clinical datasets of human stomach cancers from The Cancer Genome Atlas (TCGA)^{21,22} also reported the presence of *MYD88* gene deletions

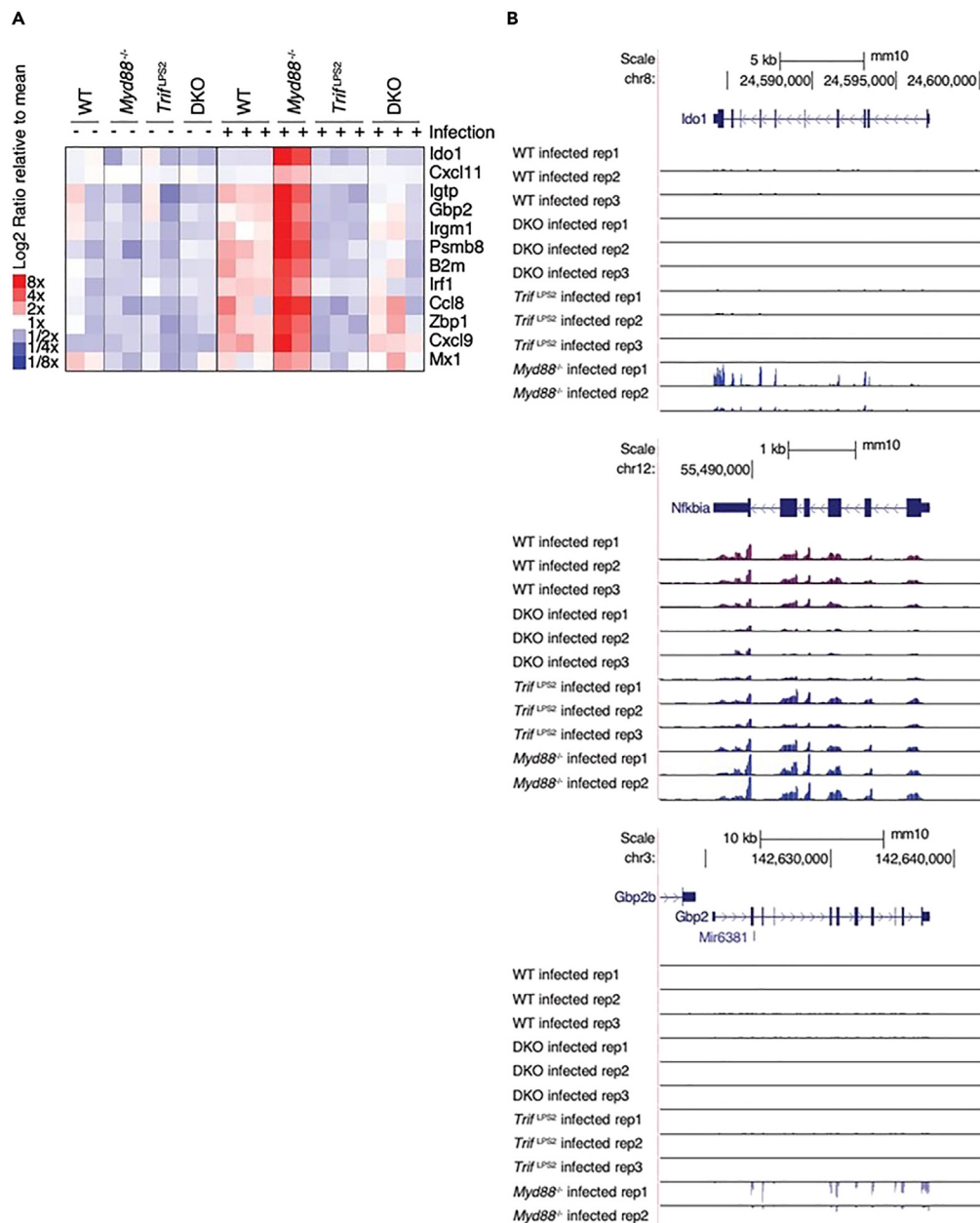


Figure 3. *Helicobacter* induced high expression of interferon-stimulated genes (ISGs) in the stomachs of mice in the absence of MyD88 signaling

(A) Relative gene expression values for several ISGs across all replicate experiments for uninfected and *H. felis*-infected gastric tissue samples from WT, *Myd88*^{-/-}, *Trif*^{ΔPS2}, and double deficient (*Myd88*^{-/-} and *Trif*^{ΔPS2}, DKO) mice.

(B) The genome tracks show normalized read density for each RNA-seq experiment at the canonical NF-κB target *Nfkbia* locus, and ISG loci *Ido1* and *Gbp2* in gastric tissue samples from WT, *Myd88*^{-/-}, *Trif*^{ΔPS2}, and double deficient (*Myd88*^{-/-} and *Trif*^{ΔPS2}, DKO) mice.

and mutations in gastric and esophageal adenocarcinomas. Considering these findings, along with data from our animal model of gastric cancer, we performed Kaplan-Meier analysis of *MYD88* expression in gastric cancer patients²³ to validate the clinical relevance of our mouse study results. The database provided data from a total of 1,065 gastric cancer patients,²³ which we divided into two groups based on their *MYD88* expression levels (high and low) based on original criteria described by Busuttill et al. 2014.²⁴ Briefly, the mean expression level of genes was used to divide patient sample into high or low expression groups with the top and bottom 33 percent of the samples defined as high or low, respectively. The resulting survival plots for gastric cancer patients indicated that those with low expression of *MYD88* had significantly low survival rates compared to patients with higher expression, as shown in overall survival, OS ($p = 0.0016$, log rank test), first progression, FP ($p = 0.0049$, log rank test), and post-progression survival, PPS ($p = 0.022$, log rank test) analyses (Figure 8). These results

Top Enriched Motifs for <i>Myd88</i> ^{-/-} vs. WT (infected)					
Motif Name	Consensus	% of Target	% of Background	-Log P-value	-Log P-value
ISRE	AGTTTCATTTTC	18.48%	1.06%	69.49	
ETS:IRF composite	MGGAAAGTGAAAC	33.15%	18.08%	14.13	
NFκB	WGGGGATTCC	14.67%	6.42%	9.792	
STAT/GAS	RTTCTNAGAAA	10.87%	4.19%	9.131	
bZIP:IRF	NAGTTTCABHTTGACTNW	8.70%	3.02%	8.66	
KLF/Sp1	DGGGYGKGGC	53.80%	41.31%	7.74	
RUNX	NWAACCACADNN	14.13%	7.24%	7.047	
TATA-box	GSCTWTAAGG	23.37%	14.54%	6.916	
Top Enriched Motifs for <i>Trif</i> ^{IPs2} vs. WT (infected)					
Motif Name	Consensus	% of Target	% of Background	-Log P-value	-Log P-value
PRDM14	RGGTCTCTAAC	12.50%	3.04%	6.48	
VDR	ARAGGTCANWGAGTTCANN	8.93%	1.79%	5.699	
T-box	AGGTGTGAAA	16.07%	6.22%	4.904	
ZNF669	GARTGGTCATCGCCC	5.36%	1.04%	3.873	
Zic	CCTGCTGAGH	21.43%	11.58%	3.69	
Nkx2.2	BTBRAGTGSN	33.93%	21.83%	3.675	
KLF/Sp1	RGGGGGCGGGC	64.29%	50.42%	3.664	
Top Enriched Motifs for DKO vs. WT (infected)					
Motif Name	Consensus	% of Target	% of Background	-Log P-value	-Log P-value
EBF	GTCCCCWGGGGA	19.94%	14.98%	4.801	
ZNF768	RHHCAGAGAGGB	1.17%	0.25%	4.487	
OCT4-SOX2-TCF-NANOG	ATTTGCATAACAATG	1.76%	0.59%	4.022	
PAX3:FKHR-fusion	ACCRTGACTAATTNN	2.05%	0.82%	3.726	
YY1	CAAGATGGCGGC	5.57%	3.38%	3.669	
STAT6	TTCKNAGAA	5.57%	3.39%	3.65	
Rbpj1	HTTCCASG	16.13%	12.69%	3.275	

Figure 4. Ranking list of significantly enriched promoter motifs

Top 8 non-redundant transcription factor motifs enriched in the promoters of genes up-regulated in gastric tissue from *H. felis*-infected *Myd88*^{-/-} mice relative to *H. felis*-infected WT, *Myd88*^{-/-}, *Trif*^{IPs2}, and double deficient (*Myd88*^{-/-} and *Trif*^{IPs2}, DKO) mice.

establish a clear link between MyD88 deficiency and gastric cancer and suggest that the level of *MYD88* expression during *Helicobacter* infection may serve as an important prognosticator of disease outcome.

We proceeded to examine the expression of *TRIF* in human gastric cancer patients using the Kaplan-Meier survival analysis.²³ The results revealed that patients with high expression of *TRIF* had a significantly lower survival rate than that of patients with lower *TRIF* expression in the OS (log rank p = 0.0077), FP (log rank p = 0.0011), and PPS (log rank p = 0.00039) analyses. This finding strongly suggests that high expression of *TRIF* is associated with poor prognosis in gastric cancer patients (Figure 9). These clinical observations align with the findings from our mouse studies, indicating that the hyperactivation of the TRIF pathway and subsequent activation of the IFN-I signaling pathway play important roles in promoting the progression of *Helicobacter*-induced gastric cancer. As a result, targeting the TRIF signaling pathway holds the potential to intercept the development of precancerous lesions, offering a potential approach to mitigate cancer progression.

DISCUSSION

Our previous study using *Myd88*^{-/-} mice suggested a potential link between TRIF activation and the progression of *Helicobacter*-induced gastric neoplasia.⁷ Upon infection with *H. felis*, these mice exhibited rapid progression of gastric pathology, including severe gland atrophy, hyperplasia, intestinal metaplasia, and dysplasia. However, the underlying molecular mechanism behind this phenotype remained unclear. In a separate study, we had also observed that *H. pylori* induced higher production of IFNα in bone marrow-derived macrophages (BMDM) from *Myd88*^{-/-} mice compared to WT mice,²⁵ indicating enhanced IFN response in the absence of MyD88 signaling. Based on these findings, we hypothesized that the downstream effects of an activated IFN-I signaling pathway, specifically the induction of ISGs, might play a role in the accelerated disease progression to preneoplastic lesions observed in *Myd88*^{-/-} mice.⁷

Recent research using a gastric MALT lymphoma model with *H. suis* infection also supported the importance of the TRIF-IFN-I signaling pathway in lymphoma development.¹¹ However, the timing of gastric epithelial changes associated with this disease progression and the role of downstream IFN-I target genes, ISGs,¹⁶ remained unknown. In the present study, we investigated these aspects and found that gastric epithelial changes occurred as early as one month after *H. felis* infection, with significant preneoplastic lesions developing at three months. Moreover, upregulation of ISGs was closely associated with severe disease pathology, including high-grade dysplasia, upon *H. felis* infection, suggesting a potential role for these ISGs in promoting the development and progression of gastric cancer.

Our *H. felis*-infected *Myd88*^{-/-} mouse model provided meaningful insights that parallel human disease. Specifically, we observed that ISGs upregulated in the mouse model were also elevated in gastric tissues from *H. pylori*-infected patients with atrophic gastritis (Figure 6).¹⁹

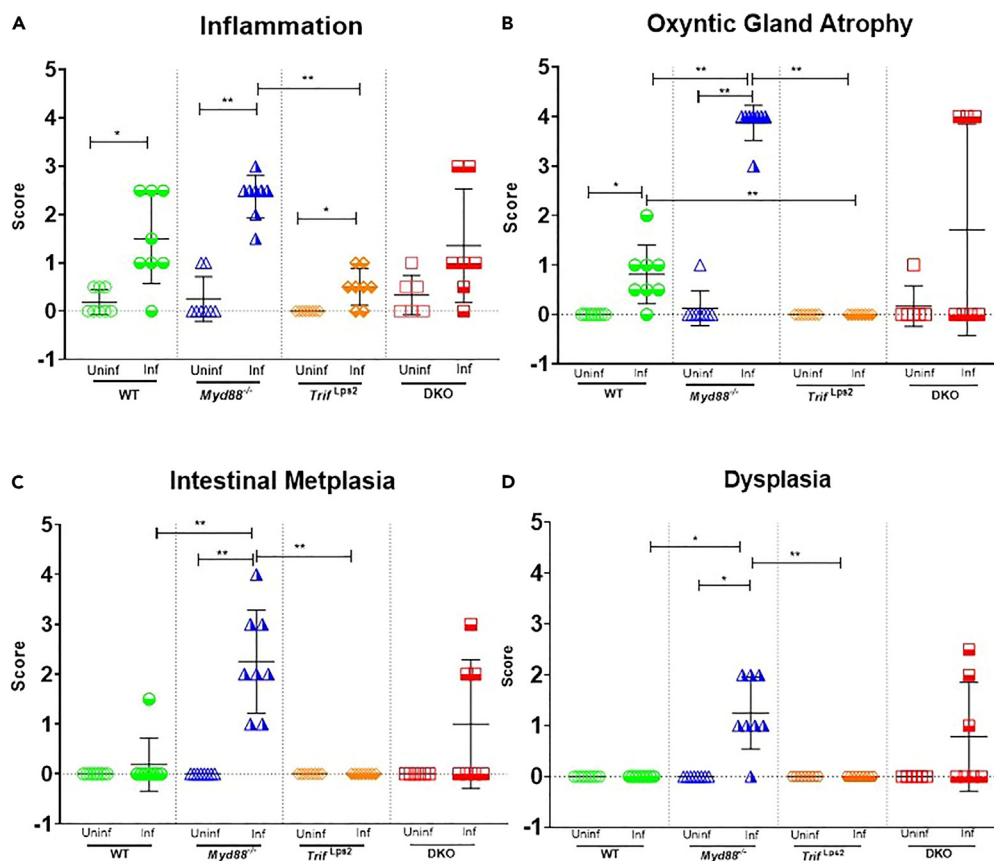


Figure 5. Histopathologic scores in mouse gastric tissue in response to *H. felis* infection

WT and *Myd88*^{-/-} *Trif*^{Lps2}, and double deficient (*Myd88*^{-/-} and *Trif*^{Lps2}, DKO) mice were infected with *H. felis* (Inf.) for 3 months (WT n = 8; *Myd88*^{-/-} n = 8; *Trif*^{Lps2} n = 8; DKO n = 7) or left uninfected (Uninf.) (WT n = 8; *Myd88*^{-/-} n = 8; *Trif*^{Lps2} n = 8; DKO n = 6). Mouse stomachs were processed and scored double-blind for histologic disease severity for inflammation (A), gland atrophy (B), intestinal metaplasia (C) and dysplasia (D) on an ascending scale from 0 (no lesions) to 4 (severe lesions). Data are represented as mean ± SEM. Data were analyzed using Mann Whitney U Test. *p < 0.01, **p < 0.001.

Moreover, our analysis of gastric biopsy samples from our UCSD biorepository showed upregulation of these ISGs in gastric cancer patients (Figure 7). Together, these data suggest that ISGs may be involved in the transition of disease toward malignancy. The upregulation of these ISGs could drive molecular changes associated with *Helicobacter*-induced disease progression, leading to severe gastric pathology and pre-cancerous lesions.

Although the role of ISGs in *Helicobacter*-induced disease pathogenesis is not well understood, some of the upregulated ISGs in mice and patient gastric tissue samples, including *Gbp2* and *Ido1*, have been associated with other human disease conditions. As an example, *GBP2*¹⁰ is highly expressed in human esophageal squamous cell carcinoma²⁶ and is associated with increased proliferation,²⁷ which aligns with the enhanced proliferation observed in our *Myd88*^{-/-} mouse infection model.⁷ *Ido1*, another highly expressed ISG in gastric tissues of *Myd88*^{-/-} mice following *H. felis* infection, is implicated in immune tolerance and high expression has been correlated with poor clinical outcomes in multiple cancer types.^{28–39} Upregulation of *ISG15* expression in colon cancer tissues has been reported to promote proliferation and metastasis and is associated with poor prognosis in colon cancer patients.⁴⁰ However, additional targeted inhibition studies are needed to directly assess the specific contribution of individual ISGs in disease pathology. Nevertheless, our results strongly indicate that upregulation of ISGs is involved in the severe gastric pathology observed in *Myd88*^{-/-} mice (Figure 1)⁷ and in human gastric cancer patients (Figures 6 and 7). Cumulatively, our data show that *Helicobacter*-induced disease progresses rapidly to severe disease pathology in the presence of an unopposed activated TRIF pathway, leading to the upregulation of ISGs. Activation of the TRIF-IFN-I-dependent signaling pathway by *Helicobacter* may create a heightened inflammatory microenvironment that disrupts mucosal gastric homeostasis, resulting in the development of a dysplastic epithelium and ultimately gastric cancer. Indeed, gastric biopsy samples from *H. pylori*-infected patients with atrophic gastritis¹⁹ showed higher expression of ISGs compared to uninfected individuals. Furthermore, high expression of *TRIF* in gastric biopsy samples from gastric cancer patients was associated with poor OS.

In conclusion, our study provides significant evidence, from both *in vivo* models and patient-derived data, indicating that the TRIF-IFN-I pathway is involved in *Helicobacter*-induced disease progression to gastric cancer. The findings highlight the central role of this pathway in gastric carcinogenesis, emphasizing the importance of further exploration in this area. Our study demonstrates the importance of examining

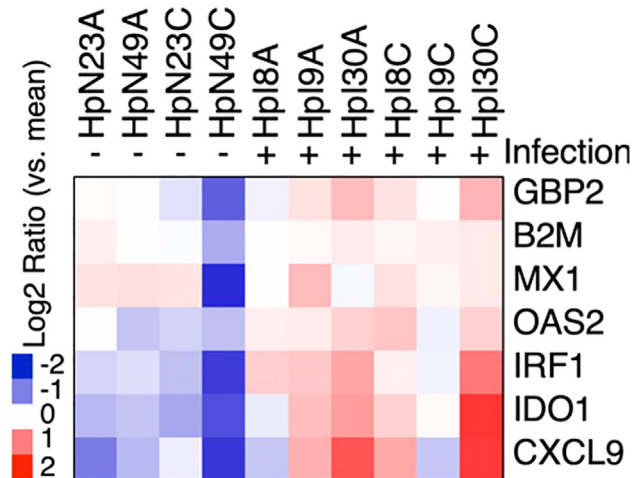


Figure 6. Heatmap showing expression of ISGs in patient gastric tissue samples

Reanalysis of published microarray dataset of gastric tissue from uninfected and *H. pylori*-infected patient with atrophic gastritis using publicly available dataset (GSE27411).¹⁹ Samples from *H. pylori*-infected patients are denoted by “+” (n = 6) and uninfected by “-” (n = 4).

the mechanisms of *Helicobacter*-induced disease during the early phase of infection. Early detection of epithelial changes could enhance the identification of drivers, leading to the discovery of effective molecular targets that may halt the progression of disease to neoplastic lesions and ultimately gastric cancer.

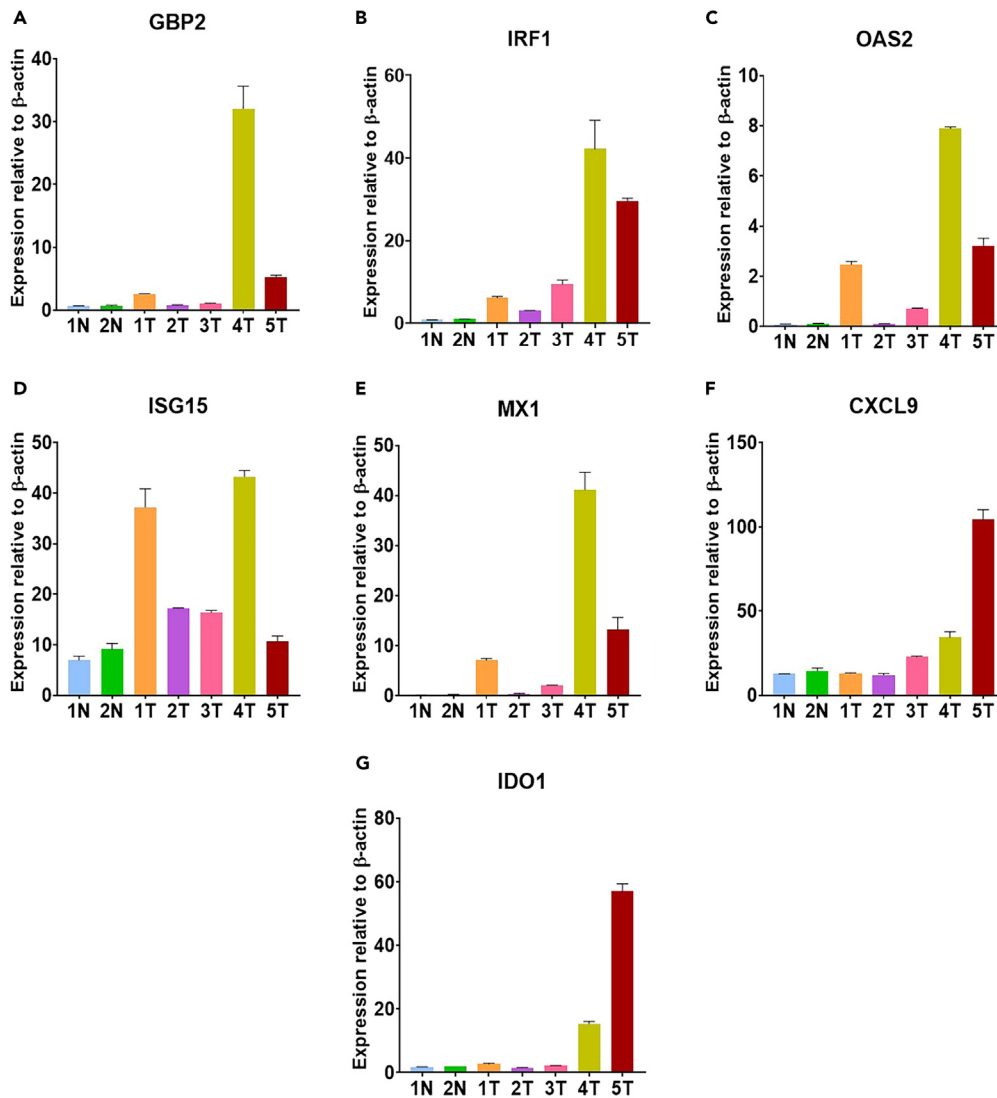
Limitations of the study

One limitation of this study is that the *H. pylori* status of gastric cancer patients from our biorepository and the Kaplan-Meier plotter is often not defined. Infection with *H. pylori* is the major cause of gastric cancer, as epidemiologic studies estimating that 75% of gastric cancer cases are a result of infection with the bacterium.^{41–43} Thus, there is a possibility that some of the gastric cancer patients in our study may not have been *H. pylori* positive. Another limitation of the study is the remaining outstanding question of whether MyD88 and TRIF interactively influence *Helicobacter*-induced gastric carcinogenesis. While we show that disease pathology in DKO mice is moderately severe, we do not yet know whether the MyD88 or TRIF pathway played a more important role. The elucidation of this point will require long-term animal studies to discriminate the key pathways and the impact of interactions between them. Further, while we used male mice to allow for comparison with previous studies that have investigated gastric carcinogenesis using the *H. felis* infection mouse model that have used only male mice,^{44–46} this limits generalization of the results. Finally, our study used *H. felis* gene and whole genome reads to estimate colonization, which is not a very quantitative way of assessing colonization. Our prior work had shown no significant differences in *H. felis* colonization between WT and *Myd88*^{-/-} mice.⁷ In the current study, where our primary focus was on disease severity rather than colonization, we found no significant differences in reads either using genes frequently used to assess colonization or whole genome reads between infected mouse strains (Figure S1).

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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1N-2N – Normal; 1T-5T – Gastric Cancer Patients

Figure 7. Expression of ISGs in patient gastric biopsies

Gastric cancer patients' gastric tissues obtained from the UCSD Biorepository showed upregulation of ISGs including *GBP2* (A), *IRF1* (B), *OAS2* (C), *ISG15* (D), *MX1* (E), *CXCL9* (F), and *IDO1* (G). N, denotes normal and T, tumor. Data are represented as mean \pm SEM.

- RNA-seq analysis
- Pathway and motif enrichment analysis
- Human gastric interferon-stimulated gene (ISG) expression
- Survival analyses of *MYD88* and *TRIF* using the Kaplan-Meier plotter
- **QUANTIFICATION AND STATISTICAL ANALYSIS**

SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.isci.2024.109457>.

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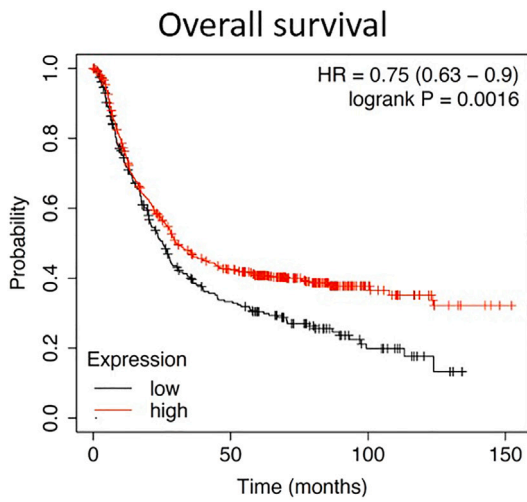
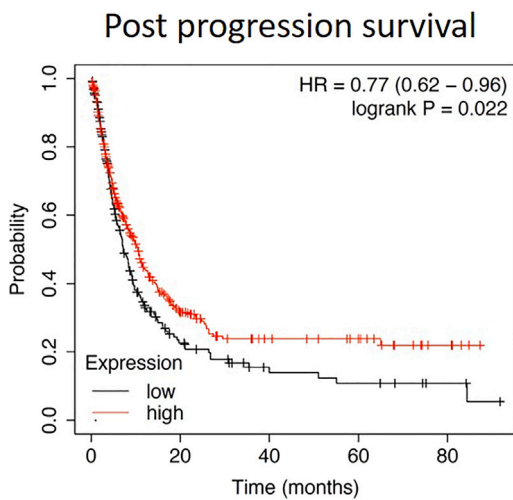
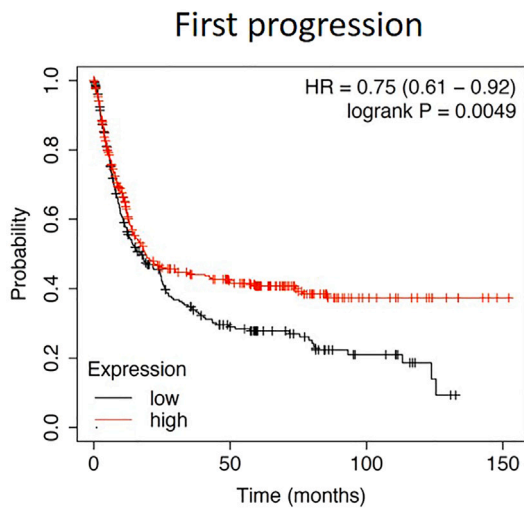


Figure 8. Kaplan-Meier survival curves for *MYD88* expression in patients with gastric cancer

This online database includes data from 1065 gastric cancer patients.²³ OS, overall survival; FP, first progression; PPS, post-progression survival.



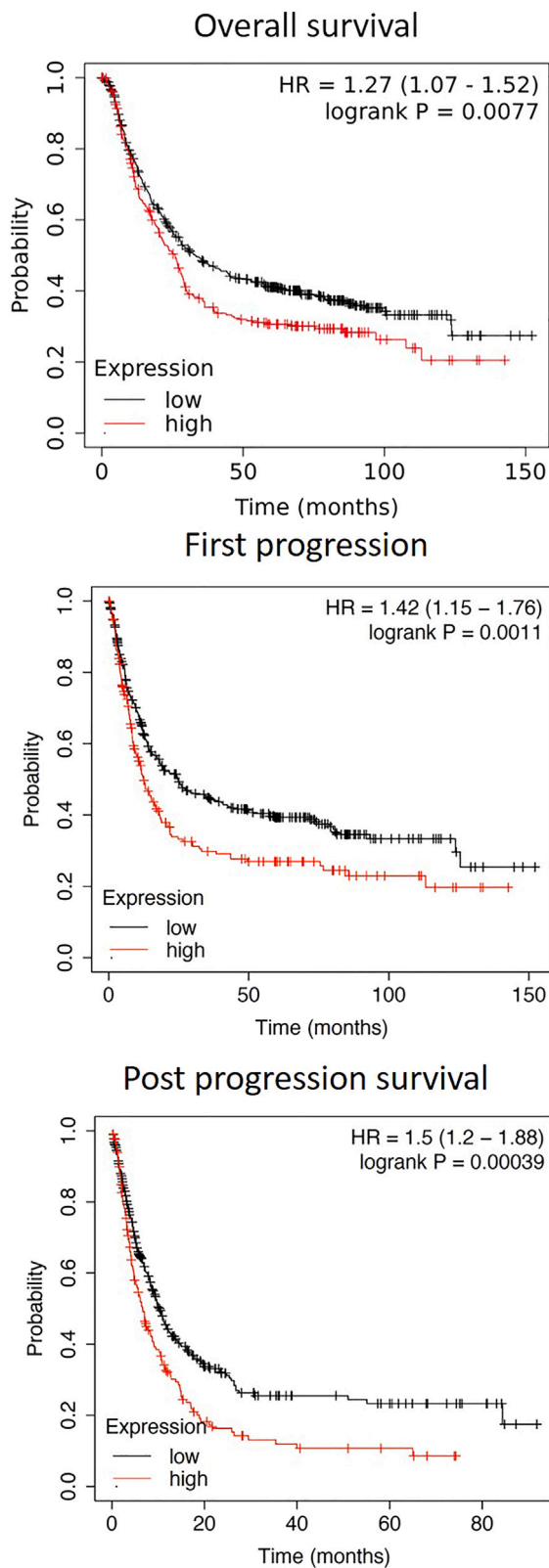


Figure 9. Kaplan-Meier survival curves for *TRIF* expression in patients with gastric cancer

This online database includes data from 1065 gastric cancer patients.²³ OS, overall survival; FP, first progression; PPS, post-progression survival.

AUTHOR CONTRIBUTIONS

M.O. supervised and designed the study concept and research; C.B. supervised data analysis; P.B., I.L.-P., and J.H. collected and analyzed data; M.V.E. performed mouse histopathologic scoring; M.C. supervised mouse breeding; M.A.T. and M.B. acquired UCSD biorepository patient information; P.B. and I.L.-P. wrote the draft of the manuscript; C.B. and M.O. revised the manuscript. All authors read and approved the manuscript.

DECLARATION OF INTERESTS

All authors declare that no conflict of interest exists.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Bacterial and virus strains		
<i>Helicobacter felis</i> , strain CS1	American Type Culture Collection	ATCC 49179
Biological samples		
Human Gastric Biopsy Tissue Samples	Biorepository, University of California, San Diego	N/A
Critical commercial assays		
Direct-zol RNA MiniPrep Kit	Zymo Research	R2050
High Capacity cDNA Reverse Transcription Kit	Thermo Fisher	4368814
Illumina TruSeq Stranded Total RNA Sample Prep Kit with Ribo-Zero	Illumina	20020597
Deposited data		
GEO accession number: GSE250438	This paper	GSE250438
Experimental models: Organisms/strains		
Mouse: WT: C57BL/6	Jackson Laboratory	C57BL/6J
Mouse: <i>Myd88</i> KO: <i>Myd88</i> deficient	Dr. Akira's laboratory	N/A
Mouse: <i>Trif</i> KO: <i>Trif</i> deficient	Dr. Akira's laboratory	N/A
Mouse: <i>Myd88/Trif</i> DKO: <i>Myd88/Trif</i> deficient	This paper	N/A
Oligonucleotides		
Primers for qPCR	see Table S2	N/A
Software and algorithms		
GraphPad Prism 8.2.0	GraphPad Software	https://www.graphpad.com/scientificsoftware/prism/
HOMER's analyzeRepeats.pl	Dr. Benner's lab	N/A
Other		
Nanodrop 2000	Thermo Fisher	ND-2000
Veriti Thermal Cycler	Applied Biosystems	4375305
StepOne	Applied Biosystems	272001471
Illumina/Solexa HiSeq 2500 sequencer	Illumina	SY-401-2501

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Marygorret Obonyo (mobonyo@health.ucsd.edu).

Materials availability

This study did not generate new unique reagents.

Data and code availability

- All data reported in this paper will be shared by the [lead contact](#) upon request.
- Single-cell RNA-seq data have been deposited at GEO and is publicly available as of the date of publication. The accession number is listed in the [key resources table](#).
- Any additional information required to reanalyze the data reported in this paper is available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS

Animals

Six- to ten- week-old wild type (WT) (n=42), MyD88 deficient (*Myd88*^{-/-}) (n=47), TRIF deficient (*Trif*^{Δ^{PS2}}) (n=46), double deficient (*Myd88*^{-/-} and *Trif*^{Δ^{PS2}}, DKO) (n=37) mice in the C57BL/6 background were used in this study. In this study male mice were used. WT mice were purchased from The Jackson Laboratory (Bar Harbor, ME). *Myd88*^{-/-} mice were from our breeding colony originally provided by Dr. Akira (Osaka University, Japan) and backcrossed 10 times onto a C57BL/6 background, bred, and maintained at our facility. The DKO mice were bred in our facility by crossing *Myd88*^{-/-} and *Trif*^{Δ^{PS2}}. All mice were housed together before and throughout the *H. felis* infection study for each genotype in a biosafety level II (BSL-2) facility with controlled temperature (23 ± 2°C) and relative humidity (45-60%) and had full access to food and water. The animal procedures were approved by the Institutional Animal Care and Use Committee at the University of California, San Diego, and conducted following accepted veterinary standards and ARRIVE guidelines.

Human participants

Seven de-identified snap-frozen human gastric biopsy samples were obtained from UCSD's Biorepository. Prior to sample collection, all patients provided written informed consent and were followed up. Details about the patients and their gastric tumor characteristics including race or ethnicity are provided in Table S1.⁴⁷ RNA was extracted from each biopsy tissue and subsequently processed for real-time PCR as described below. The study was conducted in accordance with the Declaration of Helsinki, and approved by the Institutional Review Board of the University of California, San Diego.

Bacterial growth conditions

Helicobacter felis, strain CS1 (ATCC 49179) was obtained from the American Type Culture collection (Manassas, VA). The bacteria were cultured on solid Columbia agar (Becton Dickinson, MD) supplemented with 5% laked blood under microaerophilic conditions (5% O₂, 10% CO₂, 85% N₂) at 37°C and passaged every 2–3 days, following established protocols.^{7,48–50} Prior to mouse infections, *H. felis* was cultured in liquid brain heart infusion broth (BHI, Becton Dickinson) supplemented with 10% fetal calf serum and incubated at 37°C under microaerophilic conditions for 48 h. Spiral bacteria were enumerated using a Petroff-Hausser chamber before infections.

Mouse infections

This study used a well-characterized cancer mouse model, infecting C57BL/6 mice with *H. felis* (strain CS1), a close relative of the human gastric pathogen *H. pylori*. Mice were inoculated with 10⁹ organisms in 300 μL of BHI by oral gavage using a feeding needle.^{7,48–50} Inoculation of mice with *H. felis* was repeated three times at 2-day intervals. Control mice received BHI only. At 1 month, 3 months, and 6 months post-infection, mice were euthanized, and their stomachs removed under aseptic conditions. The stomach tissue was cut longitudinally, and sections were processed for RNA extraction and histopathology. All *H. felis*-infected mice were confirmed to be colonized by assessing the expression of the flagella filament B (*flab*) gene using real-time PCR.⁷ The amplification conditions consisted of an initial cycle of 95°C for 5 min, annealing at 60°C for 20 sec, and extension at 72°C for 40 sec. Gene expression levels were normalized to glyceraldehyde 3-phosphate dehydrogenase (GAPDH). There were no significant differences in the expression of the *flab* gene among all the infected mice groups.

METHOD DETAILS

Histology

Mouse stomachs were processed for histology according to the following the procedures.^{7,51} Longitudinal sections of stomach tissue from each mouse were fixed in neutral buffered 10% formalin, embedded in paraffin, and then 5 μm sections were stained with hematoxylin and eosin (H&E). The gastric histopathologies, including inflammation, epithelial defects, gland atrophy, hyperplasia, intestinal metaplasia, and dysplasia, were evaluated and scored for histologic disease severity on a scale from 0 (no lesions) to 4 (severe lesions). The scoring was performed in a double-blind manner by a pathologist (Dr. Estrada) following the criteria outlined by Rogers et al.⁵² Specifically, the histological characteristics of the parameters were defined as follows: inflammation—filtration of inflammatory cells into the mucosa and submucosa; epithelial defects—thinned epithelial surface, epithelial and mucosal erosion and multifocal collapse; oxyntic gland atrophy—loss of oxyntic glands (which contain parietal cells and chief cells) in the gastric corpus; hyperplasia—elongation of gastric glands due to increased numbers of surface cells (foveolar cells); intestinal metaplasia—replacement of gastric epithelium by intestinal type gastric epithelium; and dysplasia—abnormal cellular and glandular maturation of the epithelium, with score of 3 indicating carcinoma *in situ*.⁵²

RNA extraction from mouse and human gastric tissue samples

RNA was extracted from gastric tissue obtained from human biopsy samples (described above) and gastric tissue samples from *H. felis*-infected and uninfected WT, *Myd88*^{-/-}, *Trif*^{Δ^{PS}}, and DKO mice. Gastric tissue samples were homogenized in 1 ml of TRIzol reagent (Invitrogen, Carlsbad, CA) following the manufacturer's instructions. RNA was extracted using the Direct-zol RNA mini kit (Zymo Research, Irvine CA) as per the manufacturer's instructions and stored at -70°C until further use. The quality of the extracted RNA was assessed using a Nanodrop

system (Thermo Fisher, Waltham MA) by measuring absorbance levels at 260 and 280 nm. Subsequently, the RNA was utilized for RNASeq analysis in the mouse gastric tissue samples and qPCR in the human gastric tissue samples.

cDNA synthesis and quantitative real-time RT-PCR

Real-time RT-PCR (reverse transcription polymerase chain reaction) was conducted the StepOne Real Time PCR system (Applied Biosystems, Carlsbad CA).^{7,10,50,51} Expression levels of specific downstream genes in the TRIF-IFN-I pathway in human gastric biopsy samples including, indoleamine 2,3-dioxygenase 1 (*IDO1*), guanylate binding protein 2 (*GBP2*), transcription factor Interferon regulatory factor 1 (*IRF1*), 2'-5'-oligoadenylate synthase 2 (*OAS2*), and chemokine (C-X-C motif) ligand 9 (*CXCL9*), MX dynamin-like GTPase 1 (*MX1*), and interferon-stimulated gene 15 (*ISG15*), were determined. RNA (2µg/sample) isolated from gastric tissue samples was reverse transcribed into cDNA using the High Capacity cDNA Reverse Transcription Kit (Thermo Fisher, Waltham MA). For amplification, 1µl of cDNA was used per well in a total of 10µl reaction mix. The amplification conditions consisted of an initial cycle of 95°C for 5 min, annealing at 60°C for 20 sec, and extension at 72°C for 40 sec. Gene expression levels were normalized to β-actin. The data collected were analyzed using comparative cycle threshold calculations (ΔCT Applied Biosystems) and plotted using GraphPad Prism software. The primers used in this study are listed in [Table S2](#).

RNA sequencing

RNA concentration and integrity were evaluated using an Agilent 2100 Bioanalyzer, and each sample was assigned an RNA integrity number (RIN). Selected samples underwent further processing into RNA-seq libraries using the Illumina TruSeq Stranded Total RNA Sample Prep Kit with Ribo-Zero (Human/Mouse/Rat), which involved ribosomal RNA depletion, fragmentation, cDNA synthesis, and cDNA library preparation. Subsequently, sequencing was performed on an Illumina/Solexa HiSeq 2500 sequencer, generating 75-bp single-end reads for all samples at a depth of approximately 40 million reads per sample.

RNA-seq analysis

For RNA-seq data analysis, the reads were aligned to the combined mouse and *H. felis* genomes (GRCm38/mm10 and NC_014810.2, respectively) using STAR with default parameters.⁵³ Only reads that aligned uniquely to a single location across genomes with a mapping quality score greater than 10 (MAPQ > 10) were used for downstream analysis. Gene expression levels were determined by counting the number of reads overlapping exons for all genes using HOMER's analyzeRepeats.pl tool and the transcriptome annotation from GENCODE (version M25).⁵⁴ Analysis was performed using DESeq2 to identify genes that were differentially expressed from unnormalized read counts.⁵⁵ Volcano plot visualization was performed using the EnhancedVolcano package from R/Bioconductor. For heatmap visualization, gene expression values were normalized using DESeq2's rlog function. Normalized genome browser read density plots were generated by creating bedGraph files with HOMER and visualizing them using the UCSC Genome Browser. Gene expression values for genes on the *H. felis* genome were determined using HOMER's analyzeRepeats.pl function based on gene annotations associated with NCBI accession number NC_014810.2, and reported as fragments per kilobase per million uniquely mapped fragments (FPKM), normalized to the total number of reads aligning to both the mouse and pathogen genomes.

Pathway and motif enrichment analysis

Pathway enrichment analysis was performed with Metascape,⁵⁶ a web-based portal that integrates multiple knowledgebases, including the Gene Ontology, to identify overrepresented pathways in sets of regulated genes. Differentially expressed genes with a fold change of at least 2 and an adjusted *p*-value of 0.05 were used as input for Metascape analysis. The pathway enrichment results reported by Metascape were corrected for multiple hypothesis testing. To analyze transcription factor motifs in mouse stomach tissue sections (RNA-seq data), we used HOMER, a software suite for DNA motif and next-generation sequence (NGS) analysis.⁵⁴ Known transcription factor motifs from the HOMER database were scanned in mouse promoter sequences defined within a range from -300 to +50 bp relative to the GENCODE-defined transcription start site (TSS) locations. The log *p*-values reported in the analysis were corrected for multiple hypothesis testing using the Benjamini–Hochberg procedure.

Human gastric interferon-stimulated gene (ISG) expression

Gene expression levels in gastric tissues were reanalyzed using data from the Gene Expression Omnibus (GEO) database, with accession number GSE27411.¹⁹ This dataset contains microarray gene expression data from gastric tissue samples obtained from individuals with and without *H. pylori* infection.¹⁹ To ensure consistency, RNA-seq expression values were quantile normalized before visualization. The resulting normalized expression values were then used to generate a heatmap using Java Tree View.⁵⁷

Survival analyses of MYD88 and TRIF using the Kaplan-Meier plotter

We utilized the Kaplan-Meier plotter bioinformatics platform (<https://kmpplot.com/analysis/>) to analyze the impact of *MYD88* and *TRIF* gene expression on overall survival (OS), first progression (FP), and post-progression survival (PPS) in gastric cancer patients. The data used for analysis was derived from the GEO database²³ and included microarray gene expression data from 1065 gastric cancer patients.



QUANTIFICATION AND STATISTICAL ANALYSIS

Statistical analysis was performed using GraphPad Prism (La Jolla, CA, USA). For normally distributed data, t-test was employed, while non-normally distributed data were analyzed using Mann Whitney *U* Test. For multiple comparisons, ANOVA with Bonferroni's correction (for normal distribution) or Kruskal-Wallis with Dunn's comparison test (for non-normal distribution) were used. Survival analysis was performed using the Kaplan-Meier method, and hazard ratio (HR) with a 95% confidence interval was calculated. Patients were divided into high and low expression groups using the best cut-off, and differences were assessed using the log-rank test. *P*-values <0.05 were considered statistically significant. Differentially expressed genes were identified as those with a fold change of at least 2 and an adjusted *p*-value of 0.05, determined by DESeq2.