

Nuclear DDX3 expression predicts poor outcome in colorectal and breast cancer

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Purpose: DEAD box protein 3 (DDX3) is an RNA helicase with oncogenic properties that shuttles between the cytoplasm and nucleus. The majority of DDX3 is found in the cytoplasm, but a subset of tumors has distinct nuclear DDX3 localization of yet unknown biological significance. This study aimed to evaluate the significance of and mechanisms behind nuclear DDX3 expression in colorectal and breast cancer.

Methods: Expression of nuclear DDX3 and the nuclear exporter chromosome region maintenance 1 (CRM1) was evaluated by immunohistochemistry in 304 colorectal and 292 breast cancer patient samples. Correlations between the subcellular localization of DDX3 and CRM1 and the difference in overall survival between patients with and without nuclear DDX3 were studied. In addition, DDX3 mutants were created for in vitro evaluation of the mechanism behind nuclear retention of DDX3.

Results: DDX3 was present in the nucleus of 35% of colorectal and 48% of breast cancer patient samples and was particularly strong in the nucleolus. Nuclear DDX3 correlated with worse overall survival in both colorectal (hazard ratio [HR] 2.34, $P < 0.001$) and breast cancer (HR 2.39, $P = 0.004$) patients. Colorectal cancers with nuclear DDX3 expression more often had cytoplasmic expression of the nuclear exporter CRM1 (relative risk 1.67, $P = 0.04$). In vitro analysis of DDX3 deletion mutants demonstrated that CRM1-mediated export was most dependent on the N-terminal nuclear export signal.

Conclusion: Overall, we conclude that nuclear DDX3 is partially CRM1-mediated and predicts worse survival in colorectal and breast cancer patients, putting it forward as a target for therapeutic intervention with DDX3 inhibitors under development in these cancer types.

Keywords: RNA helicase, DDX3X, nuclear export, CRM1

Introduction

DEAD box protein 3 (DDX3), also known as DDX3X, is a member of the DEAD box RNA helicase family of proteins. Like other family members, it has ATPase-dependent helicase activity, which allows for the unwinding of double-stranded RNA and restructuring of complex secondary RNA structures.¹ DDX3 has thus been associated with several steps of endogenous RNA metabolism, such as splicing,^{2,3} nuclear mRNA export,^{4,5} RNA interference,^{6,7} ribosomal assembly and translation initiation.^{5,8} Recent studies have also indicated a role for DDX3 in neoplastic transformation. Functional studies have shown that DDX3 has antiapoptotic properties,^{9–11} and is necessary for cell cycle progression^{12,13} and invasion.^{14,15} The tumor-enhancing role of cytoplasmic DDX3 was corroborated by studies on DDX3 expression in patient cancer samples.^{6,13}

DDX3 is known to shuttle between cytoplasm and nucleus,⁴ but in most human tissues and cell lines it is localized predominantly in the cytoplasm. We noticed that in a subgroup of colorectal and breast cancers, DDX3 can also be observed in the nucleus. It remains

to be elucidated what exact role DDX3 plays in the nucleus of cancer cells and how the subcellular localization of DDX3 is regulated in these cells. It is known that DDX3 is exported out of the nucleus as part of mRNP complexes associated both with the tip-associated protein (TAP)-dependent bulk mRNA export pathway^{5,16} and the alternative chromosome region maintenance 1 (CRM1)-dependent pathway.⁴ However, the exact nature of the relation between DDX3 expression and expression of the nuclear exporter CRM1 requires further mechanistic exploration and validation in patient samples.

Apart from the biological relevance of understanding the nuclear role of DDX3 in cancer cells, identification of nuclear DDX3 as a prognostic and therapeutic biomarker could facilitate selection of patients who would benefit from adjuvant treatment, specifically with DDX3 inhibitors that are under development.^{12,17,18} In this study we, therefore, evaluated the correlation between nuclear DDX3 expression and survival in cohorts of breast and colorectal cancer patient samples. Because DDX3 is known to bind the nuclear exporter CRM1,⁴ we sought to determine whether nuclear retention of DDX3 can be explained by aberrant CRM1 expression. Finally, we carried out *in vitro* experiments to functionally validate mechanisms of nuclear DDX3 retention.

Materials and methods

Patient samples

Tissue microarrays (TMAs) with samples from 72 colorectal cancer patients from the Academic Medical Center, Amsterdam,¹⁹ 292 colorectal cancers from Paderborn Germany¹³ and 315 breast cancers from the UMC Utrecht were used.²⁰ All TMAs included multiple cores per patient. Missing cases were due to damaged or detached cores during cutting or staining, or due to cores not containing invasive carcinoma. Clinicopathological data were retrieved from the pathology report and patient files. The breast cancer cases in this cohort were classified into molecular subtypes as were described before.^{21,22} We used anonymous archival leftover pathology material. Therefore, no ethical approval or informed consent is required according to Dutch legislation,²³ as this use of redundant tissue for research purposes is part of the standard treatment agreement with patients in our hospitals.²⁴ The medical research ethics committee of the UMC Utrecht confirmed that official approval of this study is not required (reference number WAG/mb/16/029330).

Immunohistochemistry

Four μm thick sections were cut, mounted on Super-Frost slides (Menzel & Glaeser, Brunswick, Germany),

deparaffinized in xylene and rehydrated in decreasing ethanol dilutions. Endogenous peroxidase activity was blocked with 1.5% hydrogen peroxide buffer for 15 minutes and was followed by antigen retrieval by boiling for 20 minutes in 10 mM citrate buffer (pH 6.0) for DDX3 and ethylenediaminetetraacetic acid buffer (pH 9.0) for CRM1. Slides were subsequently incubated in a humidified chamber for 1 hour with anti-DDX3 (1:1,000, pAb r647)²⁵ or anti-CRM1 (1:500, ab84375; Abcam, Cambridge, UK). After washing with phosphate-buffered saline, slides were incubated with poly-HRP-anti-mouse/rabbit/rat IgG (Brightvision; Immunologic, Duiven, the Netherlands) as a secondary antibody for 30 minutes at room temperature. Peroxidase activity was developed with diaminobenzidine and hydrogen peroxide substrate solution for 10 minutes. The slides were lightly counterstained with hematoxylin and mounted. Appropriate positive and negative controls were used throughout.

Scoring was performed by consensus of two observers (P.v.D. and M.H.v.V. or G.B.). Although the intensity of nuclear DDX3 in cells was similar, the fraction of positive cells varied. Therefore, the percentage of DDX3-positive nuclei was scored. The optimal cutoff point was selected using the online tool cutoff finder, which helps to select a cutoff that maximizes the difference in survival between groups.²⁶ Samples with $\geq 1\%$ DDX3 staining were regarded positive. Almost all cells expressed cytoplasmic DDX3, but the intensity varied and was, therefore, scored semiquantitatively as absent (0), low (1), moderate (2) or strong (3). Cases with score 0–2 were classified as having low DDX3 expression and evaluated against cases with strong expression as before.¹³ Nuclear CRM1 was scored using a semiquantitative score that was previously described by Noske et al.²⁷ The percentage of positive cells was scored as 0 (0%), 1 (<10%), 2 (10%–50%), 3 (51%–80%) or 4 (>80%). The intensity of positive nuclei was scored on the same scale as cytoplasmic DDX3. An overall score was calculated by multiplying the two scores. Cases with a score of 9 and higher were considered to have high nuclear CRM1. Cytoplasmic CRM1 was scored as absent or present.

Statistics

Clinicopathological characteristics were compared between tumors with and without nuclear DDX3. Discrete variables were compared by χ^2 or Fisher's exact test and relative risks (RRs) were calculated with 95% CI. The distribution of continuous variables was assessed graphically. Transformation was performed where indicated. Student's *t*-test and Mann–Whitney *U*-tests were calculated for normal and

non-normal distributed variables, respectively. Survival in patients with and without nuclear DDX3 was compared by plotting Kaplan–Meier curves and log-rank tests. Multivariate analysis was performed by including all factors that were found predictive ($P < 0.1$) in univariate analysis in a Cox-proportional hazard model. Stepwise backward selection based on Akaike Information Criterion was used to select the final model. Effect modifiers were identified by including multiplicative interaction terms into the model. P -values smaller than 0.05 were considered statistically significant. All statistical analyses were performed with R version 3.2.0.

Cloning and transfections

A pEGFP-C1 vector containing a parental DDX3 construct (GenBank Accession: AF061337) N-terminally fused to a 6xHis tag and EGFP sequence was used. An oligo containing three times the SV40 nuclear localization signal (NLS) sequence (5'-GATCCAAAAAAGAAGAGAAAGGTA-3', AA: DPKKKRKV) with flanking restriction sites was annealed and subcloned into the parental vector to create an NLS-EGFP-6HIS-DDX3 and an EGFP-NLS-6HIS-DDX3 construct. The DDX3 deletion constructs were created by selective amplification of the parental DDX3 construct lacking the deleted area with Phusion High-Fidelity Polymerase (New England Biosciences, Ipswich, MA, USA), followed by DpnI digestion of the bacterial backbone, gel purification, phosphorylation and religation. All constructs were verified by sequencing before usage.

HeLa cells were chosen for DDX3 localization as before.⁴ The cell line was purchased from ATCC and STR-profiled in November 2015. 2×10^4 HeLa cells were plated in a 24-well plate. After 24 hours, the cells were transfected with 250 ng DDX3 construct, 50 ng H2B-mCherry construct and 1 μ L TransIT-LT1 transfection reagent. After 12–16 hours, the cells were treated with 10 nM Leptomycin B (Sigma, St Louis, MO, USA) for 4 hours after which the number of DDX3-positive nuclei per high-power field was counted and the cells were imaged with a confocal Zeiss 780FCS system.

Results

Colorectal cancer patients with nuclear DDX3 have worse overall survival

DDX3 expression could be scored in 304 out of 364 colorectal cancers, of which 34.9% had $>1\%$ nuclear DDX3. As shown in Figure 1, the percentage of cells with nuclear DDX3 varied. Some tumors expressed nuclear DDX3 in all cells, sometimes cells with nuclear DDX3 lay dispersed among negative cells. Heterogeneity between different areas of a

tumor (different cores) was also observed. DDX3 was sometimes expressed in the nucleus of plasma cells and fibroblasts as well. Only DDX3 in the nuclei of cancer cells was considered for this analysis. As shown in Table 1, tumors with nuclear DDX3 more often had a larger tumor size ($P = 0.006$). Some tumors expressed DDX3 both in the cytoplasm and the nucleus. The presence of nuclear DDX3 correlated negatively with cytoplasmic DDX3 (RR 0.62, $P = 0.005$), which was seen in 125/304 (41%) of cases (Figure 1B).

Patients with nuclear DDX3 expression had a significantly decreased 5-year overall survival rate of 51.2%, as compared to 73.7% in those without nuclear DDX3 (hazard ratio [HR] 2.34, $P = 0.0005$; Figure 1E). Other usual predictors of survival were tumor, node, metastasis (TNM) stage ($P < 0.001$), differentiation grade ($P = 0.040$) and tumor size ($P = 0.004$). Nuclear DDX3 expression was retained in a Cox-proportional hazards model with TNM stage (HR_{adjusted} 1.69, $P = 0.057$; Table S1). No significant association between high cytoplasmic DDX3 and survival was observed in colorectal cancer patients (HR 0.69, $P = 0.160$; Figure S1).

Nuclear DDX3 is associated with worse overall survival in breast cancer patients

To assess whether our finding that nuclear DDX3 correlates with survival was also applicable to other tumors, we analyzed a cohort of 315 consecutive breast cancer cases. DDX3 expression could be evaluated in 292 breast cancer patients, of which 141 (48.3%) had nuclear DDX3 (Figure 1C). As shown in Table 2, these patients more often exhibited ductal histology ($P = 0.006$), higher grade ($P = 0.025$), larger tumor size ($P = 0.046$) and positive lymph nodes ($P = 0.003$). In addition, a trend was observed for higher mitotic activity index ($P = 0.058$). Nuclear localization of DDX3 did not correlate with cytoplasmic DDX3 expression in breast cancer patients (RR 0.91, $P = 0.493$; Figure 1D). Cytoplasmic DDX3 expression in this breast cancer cohort will be discussed more elaborately in a separate report.

As seen in Figure 1F, similar to colon cancer patients, breast cancer patients with nuclear DDX3 had a worse 5-year survival rate (75%) than those without (90%; HR 2.39, $P = 0.004$). Other variables that were significant predictors of poor survival were basal-like subtype ($P = 0.024$), positive lymph node status ($P = 0.027$), negative estrogen receptor (ER) status ($P = 0.019$), negative progesterone receptor (PR) status ($P = 0.013$) and age over 50 ($P = 0.017$). Nuclear DDX3 remained a significant predictor (HR_{adjusted} 2.63, $P = 0.010$) in a multivariate Cox-proportional hazards model with basal-like subtype, lymph node status and age (Table S2).

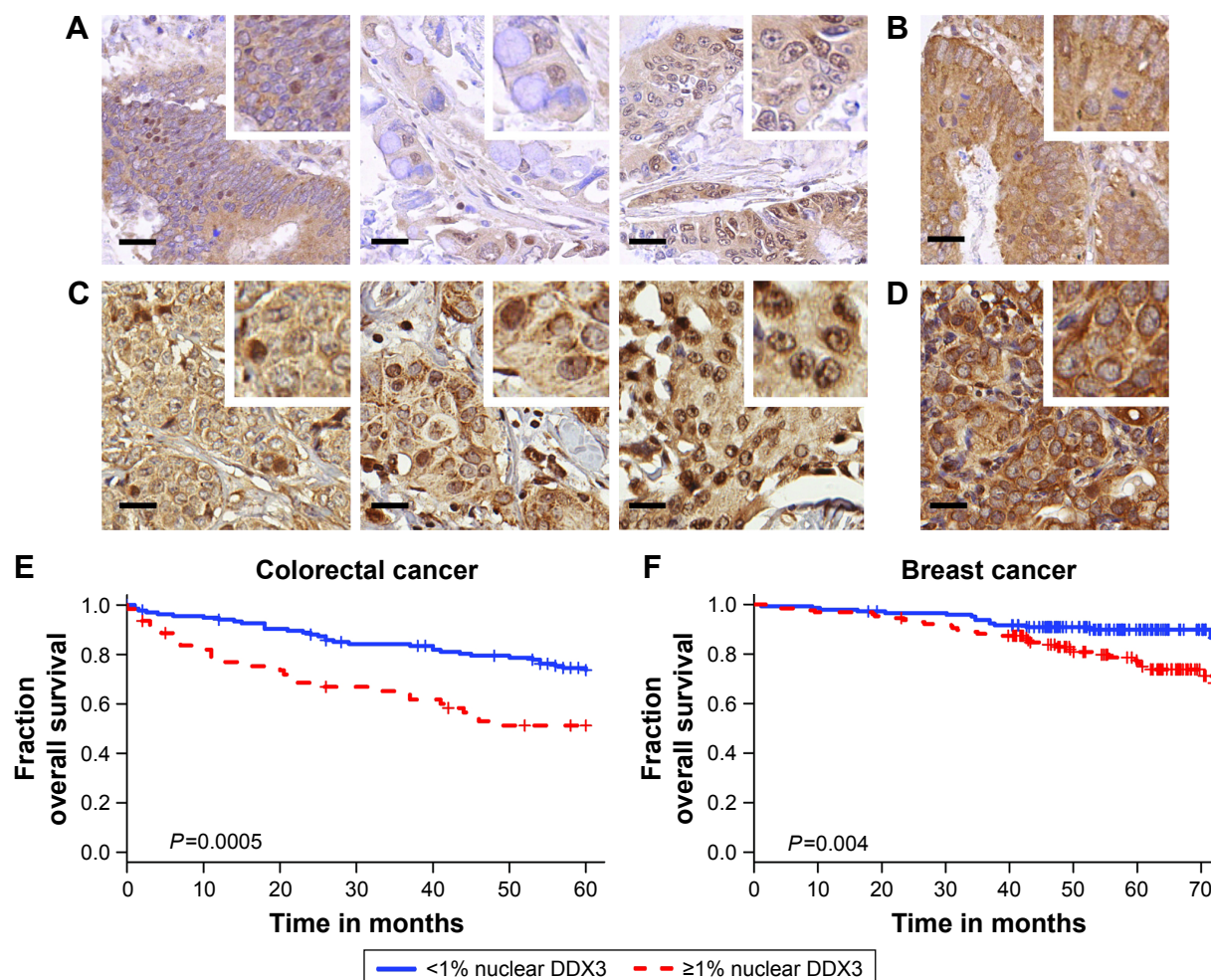


Figure 1 Nuclear DDX3 correlates with worse survival in colorectal and breast cancer patients.

Notes: Example of (A) nuclear and (B) high cytoplasmic DDX3 expression in colorectal cancer samples. Example of (C) nuclear and (D) high cytoplasmic DDX3 expression in breast cancer samples. (E) Kaplan–Meier plot showing overall survival in colorectal cancer patients with and without DDX3 expression in $\geq 1\%$ of the nuclei. (F) Kaplan–Meier plot showing overall survival in breast cancer patients with and without DDX3 expression in $\geq 1\%$ of the nuclei.

Abbreviation: DDX3, DEAD box protein 3.

The subcellular localization of the nuclear exporter CRM1 correlates with DDX3 localization

Because DDX3 is known to bind the nuclear exporter CRM1 we ascertained whether nuclear retention of DDX3 could be explained by aberrant CRM1 expression (Figure 2). CRM1 expression was mainly observed in the nucleus. High nuclear CRM1 expression was observed in 18% of colorectal cancer and 27% of breast cancer cases. Cytoplasmic expression was observed as well, more commonly in breast (36%) than colorectal cancer (8%). As shown in Figure 2A–B and Table 3, nuclear DDX3 was significantly associated with the presence of cytoplasmic CRM1 in colorectal cancers (RR 1.67, $P=0.040$). In breast cancer, no significant correlation between nuclear DDX3 and CRM1 expression was observed, suggesting that aberrant CRM1 expression could

only partly explain nuclear DDX3 retention. However, a strong correlation between the intensity of cytoplasmic DDX3 and CRM1 expression (Figure 2C and D) was observed: nuclear CRM1 was higher in tumors with high cytoplasmic DDX3 in both colorectal cancer (RR 1.77, $P<0.001$) and breast cancer cases (RR 1.75, $P=0.003$). Contrary to what was observed in colorectal cancer, a significant correlation between cytoplasmic CRM1 and high cytoplasmic DDX3 was observed in breast cancer (RR 2.45, $P<0.001$).

Nuclear DDX3 localization is due to decreased CRM1-mediated export and increased import

To characterize the functional role of DDX3 in the nucleus, we made an attempt to mimic the nuclear DDX3 expression observed in patients in an in vitro setting. We generated

Table I Correlation between nuclear DDX3 and other clinicopathological variables in colorectal cancer patients

	Total	Nuclear DDX3		RR (95% CI)	P-value
		<1%	≥1%		
Total, n (%)	304 (100)	198 (65)	106 (35)		
Sex, n (%)					0.478
Male	169 (56)	113 (57)	56 (53)	1	
Female	135 (44)	85 (43)	50 (47)	1.12 (0.82–1.52)	
Age, median (IQR)	71 (15.3)	71 (15.8)	72 (14)		0.476*
TNM-stage, n (%)					0.100
1	22 (10)	18 (12)	4 (5)	1	
2	98 (43)	67 (44)	31 (40)	1.74 (0.68–4.42)	
3	80 (35)	51 (34)	29 (37)	1.99 (0.78–5.07)	
4	30 (13)	15 (10)	15 (19)	2.75 (1.06–7.15)	
Missing	74	47	27		
Differentiation grade, n (%)					0.913
Well	16 (5)	11 (6)	5 (5)	1	
Moderate	229 (76)	149 (76)	80 (76)	1.12 (0.53–2.36)	
Poor	57 (19)	36 (18)	21 (20)	1.18 (0.53–2.63)	
Missing	2	2	0		
Tumor size (mm), mean (SD)	52 (23)	50 (23)	56 (22)		0.006**
Missing	17	12	5		
Site of origin, n (%)					0.064
Rectum	95 (31)	69 (35)	26 (25)	1	
Colon	209 (69)	129 (65)	80 (76)	1.4 (0.97–2.02)	
Histology					0.880***
Adenocarcinoma	276 (93)	179 (93)	97 (93)	1	
Mucinous	21 (7)	13 (7)	8 (8)	1.08 (0.61–1.91)	
Undifferentiated	1 (0)	1 (1)	0 (0)	NA	
Missing	6	5	1		
Cytoplasmic DDX3					0.005
Low	179 (59)	105 (53)	74 (70)	1	
High	125 (41)	93 (47)	32 (30)	0.62 (0.44–0.88)	

Notes: P-value calculated by chi-square test unless otherwise indicated. *Mann-Whitney U-test; **Student's t-test; ***Fisher's exact test.

Abbreviations: CI, confidence interval; DDX3, DEAD box protein 3; IQR, interquartile range; NA, not applicable; RR, relative risk; SD, standard deviation; TNM, tumor, node, metastasis.

a GFP-DDX3 fusion protein to study nucleocytoplasmic shuttling properties of DDX3 (Figure 3A). DDX3 is expressed primarily in the cytoplasm of HeLa cells (Figure 3B and C). Leptomycin B specifically inhibits CRM1 by covalent modification of Cys-529.^{28,29} When export of DDX3 by CRM1 was inhibited with leptomycin B, an increase in nuclear DDX3 was observed in 46% of cells, but the majority of DDX3 remained in the cytoplasm (Figure 3B and C).

Next we determined whether high nuclear DDX3 levels could potentially be explained by increased nuclear import of DDX3. We fused three tandem repeats of the SV40 NLS to the GFP-DDX3 construct (Figure 3A). Only a minor increase of nuclear DDX3 was observed in 20% of cells with the SV40 NLS, indicating that nuclear export of DDX3 is very efficient and not saturated. However, when export of the GFP-NLS-DDX3 construct was inhibited by leptomycin B, DDX3 localized strongly to the nucleus in 100% of cells, whereas cytoplasmic DDX3 was reduced to 25% of cells (Figure 3B and C). These data show that an increase in DDX3

import in combination with reduced CRM1-mediated export can result in high nuclear DDX3 expression. Experiments with an NLS-GFP-DDX3 construct yielded similar results. Unfortunately, it was not possible to create a DDX3 construct that localized to the nucleus spontaneously without inhibition of CRM1. We were, therefore, unable to study functional effects of nuclear DDX3 in an isolated fashion. However, the created constructs can be used for proof-of-principle studies on regulation of the subcellular DDX3 localization.

DDX3 localizes to the nucleolus

High expression levels of DDX3 in the nucleus allowed for closer examination of the subnuclear expression pattern of DDX3. As can be observed in Figure 3C, DDX3 is expressed diffusely within the nucleus, but is often most intense in nucleoli. Nucleolar DDX3 was seen in 2%–6% of cells after addition of either an NLS or leptomycin B. When import was increased and export was inhibited simultaneously, 30% of cells had clear localization to the nucleolus. This expression

Table 2 Correlations between nuclear DDX3 and other clinicopathological variables in breast cancer patients

	Total	Nuclear DDX3		RR (95% CI)	P-value
		<1%	≥1%		
Total, n (%)	292 (100)	151 (52)	141 (48)		
Age, median (IQR)	58 (17)	60 (18)	56 (18)		0.252*
Histology, n (%)					0.006
Ductal	231 (81)	109 (74)	122 (88)	1	
Lobular	24 (8)	19 (13)	5 (4)	0.39 (0.18–0.87)	
Medullary	31 (11)	19 (13)	12 (9)	0.73 (0.46–1.16)	
Missing	6	4	2		
Grade, n (%)					0.025
1	46 (17)	31 (22)	15 (11)	1	
2	100 (36)	54 (38)	46 (34)	1.41 (0.88–2.25)	
3	132 (48)	59 (41)	73 (55)	1.7 (1.09–2.64)	
Missing	14	7	7		
MAI, median (IQR)	13 (19)	11 (18)	15 (18)		0.058*
Missing	15	8	7		
Tumor size (mm), mean (SD)	24 (15)	23 (13)	26 (16)		0.046**
Molecular subtype					0.904***
Luminal A	224 (77)	117 (79)	107 (76)	1	
Luminal B	12 (4)	5 (3)	7 (5)	1.22 (0.74–2.00)	
HER2	10 (3)	5 (3)	5 (4)	1.05 (0.55–1.97)	
Basal-like	44 (15)	22 (15)	22 (16)	1.05 (0.76–1.45)	
Missing	2	2	0		
Lymph node status					0.003
Negative	124 (47)	73 (56)	50 (38)	1	
Positive	140 (53)	57 (44)	83 (62)	1.46 (1.13–1.88)	
Missing	29	21	8		
ER					0.927
Negative	59 (20)	30 (20)	29 (21)	1	
Positive	231 (80)	119 (80)	112 (79)	0.99 (0.74–1.32)	
Missing	2	2	0		
PR					0.840
Negative	98 (34)	51 (35)	47 (33)	1	
Positive	191 (66)	97 (66)	94 (67)	1.03 (0.80–1.32)	
Missing	3	3	0		
HER2					0.552
Negative	269 (92)	140 (93)	129 (92)	1	
Positive	22 (8)	10 (7)	12 (9)	1.14 (0.76–1.70)	
Missing	1	1	0		
Cytoplasmic DDX3					0.493
Low	194 (68)	98 (66)	96 (70)	1	
High	93 (32)	51 (34)	42 (30)	0.91 (0.70–1.19)	
Missing	5	2	3		

Notes: P-value calculated by chi-square test, unless otherwise indicated. *Mann–Whitney U-test; **Student's t-test on log-transformed data; ***Fisher's exact test.

Abbreviations: CI, confidence interval; DDX3, DEAD box protein 3; ER, estrogen receptor; Grade, Bloom and Richardson grading; IQR, interquartile range; MAI, mitotic activity index; PR, progesterone receptor; RR, relative risk; SD, standard deviation.

pattern resembled that observed in patient samples, although DDX3 expression also was strong at the nuclear membrane in those (Figure 1A and C).

Influence of CRM1 inhibition on subcellular localization of DDX3 mutants

CRM1 is known to export cargo from the nucleus by binding to a leucine-rich nuclear export signal,³⁰ which is conserved

in the first 21 amino acids of the Ded1/DDX3 subfamily of DEAD box RNA helicases.³¹ However, binding studies indicated that CRM1 binding occurs between amino acid 260 and 517 of DDX3.⁴ We evaluated whether deletion of either the N-terminal nuclear export signal (NES) domain or the 260–517 region of DDX3 results in increased retention of DDX3 in the nucleus, and whether these proteins were still responsive to CRM1 inhibition. As shown in Figure 3, deletion of the

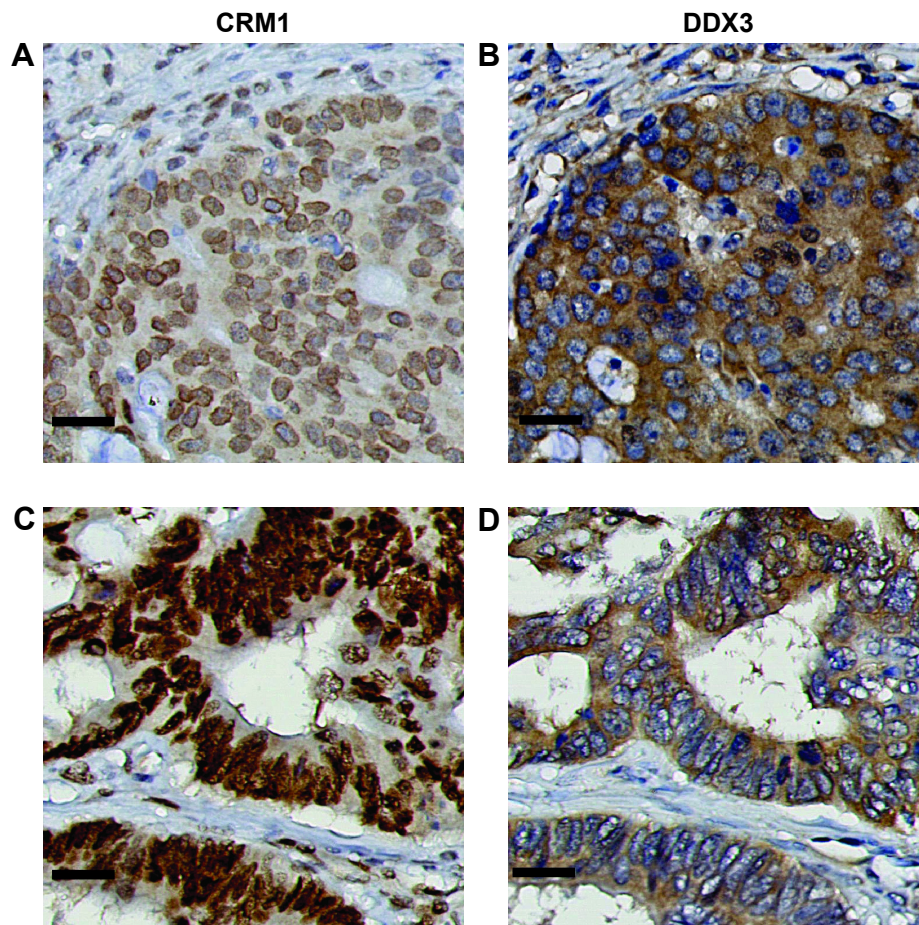


Figure 2 The subcellular localization of CRM1 correlates with the subcellular location of DDX3.

Notes: Example of how (A) cytoplasmic CRM1 expression in a colorectal cancer correlates with (B) nuclear DDX3 expression in the same tumor. Example of how (C) high nuclear CRM1 expression correlates with (D) high cytoplasmic DDX3 expression. 40× magnification. Scale bars indicate 25 μm.

Abbreviations: CRM1, chromosome region maintenance 1; DDX3, DEAD box protein 3.

Table 3 Correlation between CRM1 and nuclear DDX3 in colorectal and breast cancers

	Nuclear DDX3		P-value	Cytoplasmic DDX3		P-value
	<1%	≥1%		Low	High	
Colorectal cancer						
Cytoplasmic CRM1						
Absent	180 (95%)	87 (88%)	0.037	157 (2%)	110 (93%)	0.657
Present	10 (5%)	12 (12%)		14 (8%)	8 (7%)	
Missing	8	7		8	7	
Nuclear CRM1						
Low	156 (82%)	81 (82%)	0.952	152 (89%)	85 (72%)	<0.001
High	34 (18%)	18 (18%)		19 (11%)	33 (28%)	
Missing	8	7		8	7	
Breast cancer						
Cytoplasmic CRM1						
Absent	80 (67%)	70 (61%)	0.402	116 (75%)	33 (42%)	<0.001
Present	40 (33%)	44 (39%)		38 (25%)	45 (58%)	
Missing	31	27		40	15	
Nuclear CRM1						
Low	90 (75%)	82 (72%)	0.595	123 (80%)	48 (62%)	0.003
High	30 (25%)	32 (28%)		31 (20%)	30 (38%)	
Missing	31	27		40	15	

Note: P-value calculated by chi-square test.

Abbreviations: CRM1, chromosome region maintenance 1; DDX3, DEAD box protein 3.

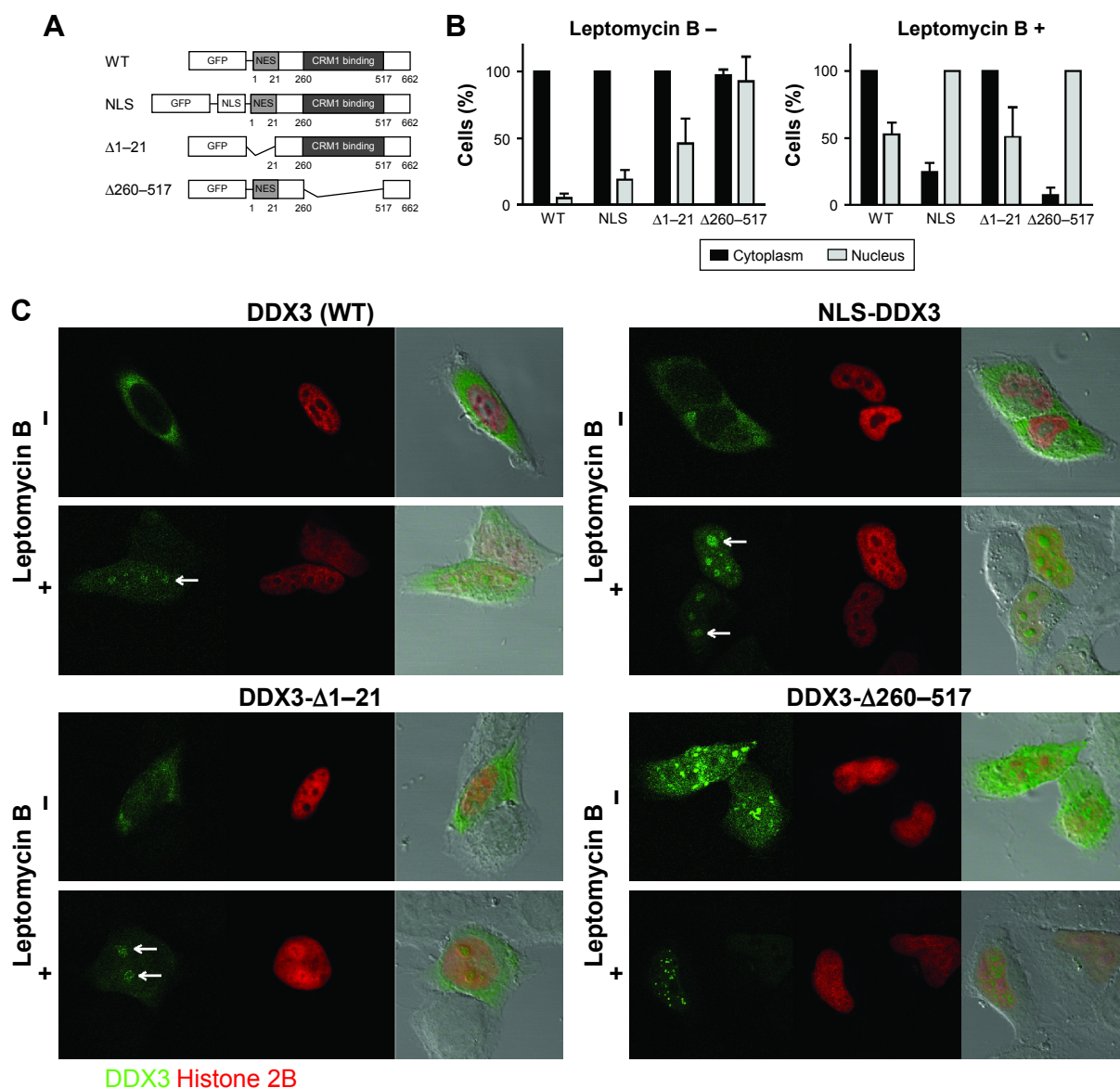


Figure 3 Influence of CRM1 inhibition on subcellular localization of DDX3 mutants.

Notes: (A) Schematic overview of the DDX3 constructs. (B) Bar graphs showing the percentage of cells with DDX3 expression in the nucleus and cytoplasm in untreated and leptomycin B-treated HeLa cells (cells can have both nuclear and cytoplasmic DDX3). Absence of error bars indicates that there was no variation, because 100% of transfected cells had DDX3 expression in that compartment. Bars represent mean percentage of positive cells of multiple microscopic fields with SD. (C) Confocal fluorescent images showing the subcellular localization of GFP-DDX3, GFP-NLS-DDX3, GFP-DDX3 $\Delta 1-21$ and GFP-DDX3 $\Delta 260-517$ (green) before and after CRM1 inhibition with 10 nM leptomycin B in HeLa cells. Nuclei are visualized by co-transfection of a Histone2B-mCherry construct (red). Nucleoli can be identified in the merged brightfield image. 40 \times magnification. Arrows indicate nucleoli.

Abbreviations: CRM1, chromosome region maintenance 1; DDX3, DEAD box protein 3; GFP, green fluorescent protein; HIV-1, human immunodeficiency virus 1; HR, hazard ratio; IQR, interquartile range; NES, nuclear export signal; NLS, nuclear localization signal; RR, relative risk; SD, standard deviation; TAP, tip-associated protein; TMAs, tissue microarrays; WT, wild-type.

first 21 amino acids containing the NES resulted in 46% of cells having DDX3 in the nucleus. No further increase was observed after leptomycin B treatment and the expression pattern was comparable to that of wild-type DDX3 after CRM1 inhibition, indicating that deletion of the N-terminal NES largely abrogates nuclear export by CRM1.

Deletion of amino acids 260–517 resulted in a speckled DDX3 nuclear expression pattern in almost all cells, but

the intensity of DDX3 remained strongest in the cytoplasm. When CRM1 was inhibited this pattern shifted to all cells having intense nuclear DDX3 expression, whereas only 7% of cells showed cytoplasmic DDX3 (Figure 3C). This indicates that although the 260–517 region has influence on the subcellular localization of DDX3, DDX3 localization is still responsive to CRM1 inhibition after deletion of this region.

Discussion

In this study we evaluated the relationship between nuclear DDX3 expression and survival in breast and colorectal cancer. We found the presence of nuclear DDX3 to be an independent predictor of worse survival in both colorectal and breast cancer. Mechanistically, in colorectal cancer nuclear DDX3 retention could in part be explained by dysregulation of the nuclear exporter CRM1, but no correlation between nuclear DDX3 and altered CRM1 expression could be observed in breast cancer. We functionally validated this finding in vitro by showing that inhibition of CRM1 with leptomycin B resulted in an increase in nuclear DDX3 levels. Analysis of the subcellular localization of DDX3 deletion mutants before and after CRM1 inhibition indicated that the N-terminal NES sequence of DDX3 is most important for this interaction. A much stronger increase in nuclear DDX3 retention was observed after addition of an NLS in combination with CRM1 inhibition, suggesting that nuclear DDX3 localization can also be regulated through nuclear import, by unknown mechanisms. Interestingly, we found DDX3 to strongly localize to the nucleolus in vitro, which resembled the expression pattern in patient samples.

DDX3 is an actively investigated molecule in cancer biology, but previous studies have focused on its cytoplasmic expression pattern in cancer cells.³² Understanding the role of nuclear DDX3 expression in tumors is relevant given DDX3's function in RNA processing and its known nucleocytoplasmic shuttling capacities. DDX3 has been found to promote oncogenesis and DDX3 inhibitors are being developed for colorectal¹³ and breast cancers¹⁷ among other malignancies.^{12,18,33,34} In addition, DDX3 is essential for the nuclear export of the human immunodeficiency virus 1 (HIV-1) protein Rev and is, therefore, also a potential therapeutic target in the treatment of HIV infections.^{35,36} Understanding the cellular mechanisms behind increased nuclear retention of DDX3 may facilitate the development of therapies specifically targeting the export function of DDX3.³⁷ In addition, nuclear DDX3 could serve as a prognostic and potentially therapeutic biomarker for selecting cancer patients that may benefit from treatment with DDX3 inhibitors.

This is the first study to describe prognostic value of DDX3 expression specifically in the nucleus. Several other studies have reported a correlation between cytoplasmic DDX3 and survival in breast and colorectal cancers.³² A study by Su et al found no significant difference in breast cancer patients with high and low cytoplasmic DDX3 expression³⁸

and reported a correlation between low cytoplasmic DDX3 expression and worse survival in colorectal cancer patients. In our larger colorectal cancer cohort, patients with high cytoplasmic DDX3 also did slightly better (HR 0.69), but this was not statistically significant. Potential explanations for the observed differences are the use of different cutoffs for positivity and different antibodies. We previously found cytoplasmic DDX3 to be associated with nuclear beta-catenin expression in patient samples and to promote oncogenic Wnt signaling. Although most colorectal cancers are driven by genetic alterations in the Wnt-signaling pathway, only a subset of cancers shows strong nuclear beta-catenin expression. Interestingly, this subgroup of colorectal cancer patients has a relatively favorable prognosis,³⁹ explaining how cytoplasmic DDX3 can be driving Wnt signaling, without being associated with worse overall survival. As DDX3 spontaneously localizes to the nucleus only in a very small percentage of cells in vitro and we were unable to create a stable cell line overexpressing full-length DDX3 in the nucleus, it is very hard to decipher the exact role of DDX3 in the nucleus on oncogenesis. However, the fact that the presence of DDX3 in the nucleus was associated with worse survival in two different cancer types indicates that cancers may benefit from high nuclear DDX3 levels as well.

When trying to understand a protein's nuclear function, it is useful to know its location in the nucleus. High DDX3 expression was specifically seen in the nucleolus. This was previously observed after HIV-1 Tat and Rev overexpression^{40,41} and DDX3 was also identified in nucleolar extracts by proteomics.⁴² The nucleolus is the structural-functional domain of the cell in which ribosomal biogenesis occurs. Prominent nucleoli have been recognized as a cytological hallmark of cancer as early as the 19th century,⁴³ but recently received renewed attention as evidence accumulates that several onco- and tumor suppressor genes are directly involved in the regulation of ribosome production to meet the altered metabolic needs of cancer cells.⁴⁴ Interestingly, DDX3 is known to play a role in ribosomal assembly and translation initiation in the cytoplasm.⁴⁵ Our observation that DDX3 localizes specifically to the nucleolus, and that this feature corresponds with worse prognosis, indicates that DDX3 may also play a role in pre-ribosomal assembly in the nucleolus. The presence of high nuclear DDX3 could reflect increased protein synthesis demands in cancers. A recent study identified essential genes in hematological malignancies and found this group to be enriched in RNA processing genes including *DDX3X*. Many of these essential genes

localized to the nucleolus.⁴⁶ Interestingly, the nucleolus is also increasingly recognized as a target for cancer therapy.⁴⁷ Further research is required to fully understand and characterize the function of DDX3 in this subcellular compartment.

With regard to the mechanism behind nuclear DDX3 retention, we observed a correlation between cytoplasmic expression of CRM1 and nuclear DDX3 expression in colorectal cancer. This suggests that dysregulation of CRM1 is one of the mechanisms of nuclear DDX3 expression. The binding site of DDX3 to CRM1 is a matter of debate. DDX3 has a classical N-terminal NES sequence that is conserved and required for CRM1 binding in the *Saccharomyces cerevisiae* homolog Ded1p^{48,49} and the *Xenopus laevis* homolog An3.³¹ However, Yedavalli et al observed binding between CRM1 and amino acids 260–517 of DDX3.⁴ Our analysis of DDX3 deletion mutants showed that deletion of both areas resulted in an increase in nuclear DDX3, but only the construct that lacked the NES lost its sensitivity to CRM1 inhibition, indicating that this is the essential domain for CRM1-mediated export. A similar conclusion was recently made by Fröhlich et al, who found the N-terminal to be essential for DDX3 transportation out of the nucleus into cytoplasmic unspliced HIV-1 mRNA ribonucleoprotein complexes.⁵⁰ It is possible that the 260–517 region of DDX3 is necessary for binding other exporters of DDX3 like TAP.⁵

As cytoplasmic CRM1 expression was infrequent in colorectal cancer and did not correlate with nuclear DDX3 in breast cancer, dysregulation of CRM1 is not likely the sole mechanism behind nuclear DDX3. When, in addition to nuclear export inhibition, we stimulated nuclear import by addition of an NLS, we observed a complete shift of all DDX3 in the cell to the nucleus, showing that increased import can also contribute to increased nuclear DDX3 levels. However, the mechanism behind nuclear import of DDX3 remains unknown. DDX3 has a classical NLS sequence at amino acid 212,⁵¹ but it is also possible that DDX3 enters the nucleus as part of a complex. Future research on this topic is warranted.

Conclusion

Nuclear DDX3 expression predicts worse survival in breast and colorectal cancer. Mechanistically, this can be partly explained by altered expression of the nuclear exporter CRM1.

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Disclosure

Venu Raman, Guus M Bol and Paul J van Diest have received a patent for the use of DDX3 as a cancer biomarker (US9322831). Venu Raman and Paul J van Diest are on the advisory board of Natsar Pharmaceuticals. The authors report no other conflicts of interest in this work.

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Supplementary materials

Table S1 Univariate and multivariate Cox-proportional hazard model of survival in colorectal cancer patients

	Univariate		Multivariate	
	HR (95% CI)	P-value*	HR (95% CI)	P-value**
Nuclear DDX3				
<1%	1		1	
≥1%	2.34 (1.43–3.85)	<0.001	1.69 (0.98–2.90)	0.057
TNM stage				
I	1		1	
2	2.9 (0.38–21.98)		2.56 (0.34–19.48)	0.364
3	7.82 (1.06–57.70)		7.15 (0.97–52.86)	0.054
4	44.34 (5.89–333.74)	<0.001	34.01 (4.45–260.15)	<0.001
Differentiation grade				
Well	1			ns
Moderate	1.24 (0.38–4.00)			
Poor	2.38 (0.71–8.00)	0.041		
Tumor size				
<40 mm	1			ns
40–60 mm	2.68 (1.28–5.63)			
>60 mm	3.57 (1.61–7.89)	0.004		

Notes: All variables significantly associated ($P < 0.1$) in univariate analysis were entered into the multivariate Cox-proportional hazards model. *P-value calculated by log-rank test. **P-value of coefficient.

Abbreviations: CI, confidence interval; DDX3, DEAD box protein 3; HR, hazard ratio; ns, no significant change in Akaike Information Criterion observed by stepwise backward selection and therefore not included in the final multivariate model; TNM, tumor, node, metastasis.

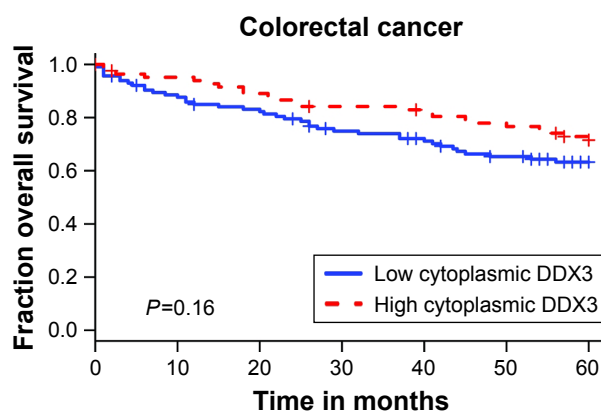


Figure S1 Kaplan-Meier plot depicting overall survival in high and low DDX3-expressing colorectal cancer samples.

Abbreviation: DDX3, DEAD box protein 3.

Table S2 Univariate and multivariate Cox-proportional hazard model of survival in breast cancer patients

	Univariate		Multivariate	
	HR (95% CI)	P-value*	HR (95% CI)	P-value**
Nuclear DDX3				
< 1%	1	0.004	1	0.01
≥ 1%	2.39 (1.29–4.43)		2.63 (1.26–5.51)	
MAI				
< 12	1	0.071		ns
≥ 12	1.78 (0.94–3.34)			
Molecular subtype				
Non-basal-like	1	0.024	1	0.045
Basal-like	2.27 (1.18–4.40)		2.17 (1.02–4.61)	
Lymph node status				
Negative	1	0.027	1	0.048
Positive	2.17 (1.08–4.39)		2.06 (1.01–4.23)	
ER				
Negative	1	0.019		
Positive	0.48 (0.26–0.90)			
PR				
Negative	1	0.013		
Positive	0.49 (0.27–0.87)			
Age				
< 50	1	0.017	1	0.017
≥ 50	2.94 (1.16–7.45)		3.55 (1.26–10.06)	

Notes: All variables significantly associated ($P < 0.1$) in univariate analysis were entered into the Cox-proportional hazards model, except for ER and PR receptor status, because these are included in the molecular subtype algorithm. *P-value calculated by log-rank test. **P-value of coefficient.

Abbreviations: CI, confidence interval; DDX3, DEAD box protein 3; ER, estrogen receptor; HR, hazard ratio; MAI, mitotic activity index; ns, no significant change in Akaike Information Criterion observed by stepwise backward selection and therefore not included in the final multivariate model; PR, progesterone receptor.

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