ARTICLE
Cellular and Molecular Biology

Statmin levels alter PTPN14 expression and impact neuroblastoma cell migration

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BACKGROUND: Statmin mediates cell migration and invasion in vitro, and metastasis in vivo. To investigate statmin’s role on the metastatic process, we performed integrated mRNA–miRNA expression analysis to identify pathways regulated by statmin.

METHODS: miRNA and gene arrays followed by miRNA-target-gene integration were performed on statmin-depleted neuroblastoma cells (CtrlmiRNA vs. Stmn Seq2miRNA). The expression of the predicted target PTPN14 was evaluated by RT-qPCR, western blot and immunohistochemistry. Gene-silencing technology was used to assess the role of PTPN14 on proliferation, migration, invasion and signalling pathway.

RESULTS: Statmin levels modulated the expression of genes and miRNA in neuroblastoma cells, leading to a deregulation of migration and invasion pathways. Consistent with gene array data, PTPN14 mRNA and protein expression were downregulated in statmin-depleted neuroblastoma cells and xenografts. In two independent neuroblastoma cells, suppression of PTPN14 expression led to an increase in cell migration and invasion. PTPN14 and statmin expression did not act in a feedback regulatory loop in PTPN14-depleted cells, suggesting a complex interplay of signalling pathways. The effect of PTPN14 on YAP pathway activation was cell-type dependent.

CONCLUSIONS: Our findings demonstrate that statmin levels can regulate PTPN14 expression, which can modulate neuroblastoma cell migration and invasion.

British Journal of Cancer (2020) 122:434–444; https://doi.org/10.1038/s41416-019-0669-1

BACKGROUND
Neuroblastoma is the most common paediatric solid tumour responsible for 15% of all paediatric oncology deaths.1  Widely disseminated, metastatic disease at diagnosis is very common and confers a poor prognosis with a 5-year survival rate of <50%.2 There is an urgent need to better understand metastasis mechanism and to identify key regulators of metastasis in high-risk neuroblastoma.

Metastasis is the movement of cancer cells from the primary sites via blood vessels and regrowth at distant sites through a highly selective process consisting of a series of discrete and sequential steps modelled into a metastatic cascade.3 Despite the focus on the actin cytoskeleton on cancer metastasis, there is increasing evidence that microtubules and their interacting proteins are also involved in this process.4 The microtubule-destabilising protein statmin is a small cytosolic phosphoprotein that is overexpressed in many cancers including neuroblastoma.5–8 We have previously shown that statmin can mediate neuroblastoma cell migration, invasion and transendothelial migration in both in vitro and in vivo neuroblastoma metastasis models.9,10 Our studies have also provided the first evidence that statmin’s influence on neuroblastoma cell migration is in part mediated by RhoA/ROCK signalling in a microtubule-independent manner.10 However, how statmin influences neuroblastoma metastasis has not yet been comprehensively addressed in neuroblastoma.

Growing evidence has revealed that microRNAs (miRNAs) play a role in multiple pathways in cancer and actively contribute to tumour development and progression.11 Specific miRNAs, collectively termed ‘metastamirs’, have multiple regulatory functions in different steps of the metastatic cascade.12 Recent work in neuroblastoma suggests that the deregulation of miRNAs may play an important role in pathogenesis and chemotherapy resistance.13 In this study, we hypothesised that new insights into the role of statmin in metastatic neuroblastoma can be gained by identifying deregulated miRNAs and associated signalling pathways following the alteration of statmin expression.

We performed functional mRNA and miRNA expression profiling in statmin-depleted neuroblastoma cells and identified differentially expressed miRNAs and their target genes. We focused on

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miRNA and gene expression profiling

Total RNA was isolated by using the miRNeasy kit (Qiagen, Victoria, Australia) according to the manufacturer’s recommendations. RNA concentration and purity were determined by absorbance using the Nanodrop 1000 (Thermo Fisher Scientific, MA, USA). RNA integrity was verified by means of ribosomal RNAs 18S and 28S on a total RNA Nano chip using the Bioanalyzer (Agilent, CA, USA). Only samples with a RNA integrity number (RIN) greater than 9 were used for array analysis. Labelling and hybridisation to Affymetrix Human 2.0 Gene ST arrays for gene expression profiling, and to Affymetrix GeneChip miRNA 3.0 arrays (based on miRBase v17) for miRNA expression profiling, was performed by the Ramacotti Centre for Genomics (University of New South Wales, Sydney, Australia). Three independent samples were prepared from CtrlshRNA and Stmn Seq2shRNA, SK-N-BE(2)/TGL cells and used for hybridisation on separate array chips.

miRNA and gene expression analysis

Analysis and visualisation of microarray data for genes and miRNAs were conducted using Partek® Genomics Suite TM software (Partek Inc., MO, USA). Background correction, quantile normalisation, log2 transformation and probe set summarisation were performed using default settings for the Robust Multichip Average (RMA) procedure. Quality of chips was assessed using QC metrics. Multidimensional intensity data were explored for differences in samples using principal component analysis (PCA). Differential expression between miRNAs or miRNAs in stathmin-depleted Stmn Seq2shRNA compared with CtrlshRNA cells was compared by using one-way analysis of variance (ANOVA) with p-value < 0.05. Significantly deregulated miRNAs or miRNAs in stathmin-depleted Stmn Seq2shRNA compared with CtrlshRNA cells were also identified based on the criteria of fold-change FC < −1.4 or FC > 1.4. Multiple comparison correction was performed using a FDR (false discovery rate) of 0.05. Selection of miRNAs and genes, which were differentially expressed in Stmn Seq2shRNA compared with CtrlshRNA, was illustrated using volcano plots. Unsupervised hierarchical clustering was employed for visualisation of patterns in the data. Intensity values were therefore standardised to a mean of zero and scaled to a standard deviation of one. Agglomerative hierarchical clustering of significantly deregulated miRNAs or miRNAs was performed using Euclidean distance to determine row/column dissimilarities, while the distance between two clusters was computed using average linkage. Dendrograms were used to visualise the hierarchy of clusters and identify samples and miRNAs or genes with similar profiles.

Gene enrichment analysis

Gene set enrichment analysis (GSEA) and Gene Ontology (GO) analysis were performed to identify cellular processes and function in which genes with significantly differential expression are involved in. Enrichment of gene sets and GO analysis was determined based on gene enrichment scores and p-values, which indicate if the differentially expressed genes belong to a certain category more often than expected randomly. The GO enrichment analysis indicated if certain GO terms were overrepresented in the list of differentially expressed genes. Enrichment scores and p-values indicate if the differentially expressed genes belong to a certain category more often than expected randomly. GO terms with enrichment p-values < 0.05 were selected as significant.

Integration of microarray data by correlating differentially expressed miRNAs and target genes

The integration of miRNA and mRNA expression data was performed using a tool integrated in the Partek software (Partek Inc.) and based on miRNA-target-gene databases. Multiple integration analyses were performed with the commonly used target prediction databases Targetscan (version 6.2) (www.TargetScan.org) and Microcosm (version 5) (https://omictools.com/microcosm-targets-tool), and the results were intersected. Only miRNAs and mRNAs, which were previously identified as significant or differentially expressed between CtrlshRNA and Stmn Seq2shRNA, were included in the analysis.

RNA isolation and real-time quantitative PCR

Total RNA was extracted using the RNeasy kit (Qiagen) and reverse transcribed using either the High Capacity cDNA Reverse Transcription kit (ABI, CA, USA) or QuantiTect Reverse Transcription kit (Qiagen) for gene expression analysis. RT-qPCR was performed using a Power SYBR green PCR master mix (Applied Biosystems, CA, USA) with STMN1_1_SG QuantiTect Primer assay (Qiagen), PTPN14 primers (Geneworks) or CYR61_1_SG QuantiTect Primer assay (Qiagen) using an Applied Biosystems 7900HT Fast Real-time PCR System. Stathmin and PTPN14 expression were normalised to the housekeeping gene, β2-microglobulin (QuantiTect primer assay, Qiagen).

For miRNA expression analysis, 500 ng of total RNA was reverse transcribed using the Taqman microRNA Reverse Transcription kit (Applied Biosystems) and the Taqman microRNA assays (Applied Biosystems). Preamplification of the reverse transcription product was performed using the Taqman Preamp master mix (Applied Biosystems) according to the manufacturer’s instructions. RT-qPCR was performed using the TaqMan® Universal PCR Master Mix, no AmpErase® UNG (Applied Biosystems) with miR132, miR221, miR222, miR-382, miR-488, miR620, miR-935, miR1281, miR3935, miR-4656, miR4492 and miR4682 Taqman microRNA assays (Applied Biosystems) using an Applied Biosystems 7900HT Fast Real-time PCR System (Applied Biosystems). Expression of miRNA was normalised to the controls RNU48, RNU6B and RNU58A (Taqman microRNA control assays, Applied Biosystems).

Transfection assays

Target-gene expression was determined in SK-N-BE(2) cells transfected with miR/Vana miR-382-5p mimic (1 pm) (Ambion, Thermo Fisher Scientific) or miR/Vana miRNA Mimic Negative control #1 (1 pm) (Ambion, Thermo Fisher Scientific), as well as miR/Vana miR-382-5p inhibitor (100 nM) (Ambion) or miR/Vana miRNA Inhibitor Negative control #1 (100 nM) (Ambion) using
Lipofectamine 2000 (Life Technologies, Victoria, Australia) according to the manufacturer’s instructions. Cells were harvested 48 h post transfection.

Transfection with siRNAs was performed as previously described. Briefly, SK-N-BE(2)/TGL and SH-SYSY/TGL cells were transfected with PTPN14 siRNA sequence 3 (PTPN14 Seq3, 5′-GCUAUGACCUUUGCUU-3′), sequence 4 (PTPN14 Seq4, 5′-GGUGACCAUCUGCGAAA-3′) (5 nM) (Dharmacon, CO, USA) or AllStar control siRNA (Qiagen) using Lipofectamine 2000 (Life Technologies) according to the manufacturer’s instructions.

Western blot analysis
Protein extraction was carried out as previously described. Fractionation of nuclear and cytoplasmic protein was carried out using the NE-PER™ nuclear and cytoplasmic fraction reagent (Thermo Fisher Scientific) according to the manufacturer’s instructions. Fifteen micrograms of nuclear protein lysates and equivalent amount of the cytoplasmic protein lysates were separated on 4–15% Precast PAGE gels (Bio-Rad) and electro-transferred to nitrocellulose membranes. The membranes were probed with antibodies against PTPN14 (Sigma-Aldrich, NSW, Australia), YAP (clone D8H1X, Cell Signaling, MA, USA), Taz (clone V386, Cell Signaling), stathmin (BD Bioscience, Victoria, Australia), Topoisomerase 1 (Novus Biologicals, CO, USA) as a control for nuclear fraction and GAPDH (clone 6C5, Abcam, Victoria, Australia) as a control for equal loading. Proteins were detected by ECL Plus (Pierce, Thermo Fisher Scientific) and membranes were either scanned using the Typhoon (GE Healthcare) or exposed to the film.

Immunohistochemistry
Animal experiments were approved by the Animal Ethics Committee, University of New South Wales (ACEC #13/116B). Tumour sections were obtained from control and stathmin-depleted cells xenograft previously obtained in our lab. Previously generated shRNA-expressing SK-N-BE(2)/TGL cells (control vs. stathmin) were injected via the lateral tail vein into 6- to 8-week-old male severe combined immunodeficiency (SCID)-Beige mice. Animals were housed with an inverse 12-h day–night cycle with lights on at 8:30 pm in a temperature- (22 ± 1 °C) and humidity- (55 ± 5%) controlled room. All mice were allowed free access to water and a maintenance diet. For in vivo experiments, at least 10 mice per group were shown to have 88.5% power to detect a 50% decrease in tumour growth and/or metastases in mice implanted with stathmin shRNA neuroblastoma cells (α = 0.05). Mice were humanely killed, and organs and tumours collected at 23 days post neuroblastoma cell injection or earlier if the mice were of ill health. Immunohistochemistry was carried out as previously described. Tumour sections obtained from control and stathmin-depleted cells xenograft were incubated with PTPN14 antibody or rabbit ImmunoglobulinG control antibody at the same concentration followed by incubation with 3,3′-Diaminobenzidine as a substrate for the peroxidase reaction and haematoxylin as the counterstain. Four individual mouse samples per condition were viewed and imaged using a x60 objective in an Olympus microscope and analysed using the cellSens software.

Migration and invasion assays
Following PTPN14 or control siRNA 72 h post transfection, migration and invasion assays were carried out as previously described.

Cell proliferation assay
The PTPN14 or control siRNA-transfected cells were harvested, and total cells were counted by trypan blue exclusion at 24, 48 and 72 h. Analysis of cell growth was carried out as previously described.

RESULTS

Differential expression of stathmin modulates both gene and miRNA expression in SK-N-BE(2)/TGL neuroblastoma cells

We have previously shown that stathmin levels are important for the migration and invasion phenotype of neuroblastoma. To further investigate the role of stathmin expression in neuroblastoma metastasis, we used SK-N-BE(2)/TGL neuroblastoma control cells (CtrlshRNA) and their respective stable-knockdown stathmin cells (Stmn Seq2shRNA) (Supplementary Fig. 1a). mRNA and miRNA expression levels in Stmn Seq2shRNA and CtrlshRNA cells were measured using microarrays. This was followed by the identification of the predicted target genes of these differentially expressed miRNAs (Supplementary Fig. 1b). From microarray analysis, the fold change (FC) in stathmin expression between the CtrlshRNA and Stmn Seq2shRNA cells was FC = 1.4. Thus, miRNAs were considered differentially expressed between CtrlshRNA and Stmn Seq2shRNA cells when FC < −1.4 or FC > 1.4, and p < 0.05 (ANOVA). To get a first indication of the difference in miRNA expression between the samples, principal component analysis (PCA) was conducted. PCA showed that samples derived from CtrlshRNA and Stmn Seq2shRNA could be clearly spatially separated into two groups based on their intensity values, indicating differences in gene expression between the samples (Fig. 1a). As visualised using a volcano plot (Supplementary Fig. 2a), 324 genes were identified to be differentially expressed between Stmn Seq2shRNA and CtrlshRNA cells. This selection of differentially expressed genes included 140 up- and 184 downregulated miRNAs (Supplementary Table 1). To illustrate the difference in miRNA expression, two-way hierarchical clustering was applied to the two sets of data and the 324 deregulated miRNAs. The resulting dendrogram showed that Stmn Seq2shRNA versus CtrlshRNA Samples were clearly separated into two distinct groups based on the miRNA expression profile (Fig. 1b). In the next step, we performed microarray analysis on miRNAs derived from the two cell lines. Like the gene expression data, PCA of miRNA expression (measured as the intensity values derived from miRNA arrays) showed spatial separation between CtrlshRNA and Stmn Seq2shRNA (Fig. 1c), suggesting that reducing stathmin expression led to alterations in miRNA expression. The criteria for differential expression of miRNAs were as previously defined for mRNA expression (p-value < 0.05 and FC < −1.4 or FC > 1.4). Thirty-six miRNAs met these criteria and were therefore considered to be differentially expressed in Stmn Seq2shRNA compared with CtrlshRNA (Supplementary Fig. 2b). Out of these miRNAs, 23 were upregulated and 13 were downregulated (Supplementary Tables 2, 3). Hierarchical clustering produced a similar dendrogram as observed for the miRNA expression analysis, demonstrating the differences in miRNA profiles between the two sample groups (Fig. 1d).

Integrated approach to determine miRNAs and their predicted target genes significantly modulated in stathmin-depleted cells

To identify potential targets of miRNAs that are differentially expressed between Stmn Seq2shRNA and CtrlshRNA, we used the miRNA target-gene prediction tools, TargetScan and microCosm. Initially, 13 miRNAs and 40 target genes that were either positively or negatively correlated using Pearson’s correlation were identified. Because miRNAs downregulate expression of their targets, we focussed on 12 miRNAs and 20 target genes, which were inversely correlated in their expression. Five of these miRNAs were...
significantly upregulated, while their target genes were significantly downregulated, and seven miRNAs were significantly downregulated, while reciprocally their target genes were upregulated (Table 1).

Modulation of stathmin expression leads to deregulation of migration and invasion pathways

To determine the biological and functional significance of the differentially expressed genes, we performed GSEA and GO
Fig. 1 Differential expression of stathmin modulates both gene and miRNA expression in SK-N-BE(2)/TGL neuroblastoma cells. a Scatter plot of principal components computed from miRNA expression data exploring high-dimensional data for similarities and dissimilarities between samples. Each of the points in the plot represents a sample (chip). A code colour was used to denote samples derived from CtrlshRNA (red) and Stmn Seq2shRNA (blue). b Hierarchical clustering of 324 differentially expressed genes in Stmn Seq2shRNA versus CtrlshRNA SK-N-BE(2)/TGL cells. Genes are represented in columns and samples are shown in rows. Using average linkage and agglomerative clustering samples and genes with similar profiles were clustered. The resulting dendrogram shows a clear separation of CtrlshRNA versus Stmn Seq2shRNA samples based on the signature of the 324 differentially expressed genes and allowed for visualisation of genes, which are up-/downregulated in Stmn Seq2shRNA compared with CtrlshRNA cells. Expression was standardised to a mean of zero and a standard deviation of one. Colours are indicative of gene expression: downregulated genes have negative values and are coloured in green, while upregulated genes have positive values and are coloured in red. c Scatter plot of principal components computed from miRNA expression data. Same analysis is used as in a. d Hierarchical clustering of 36 differentially expressed miRNAs in Stmn Seq2shRNA versus CtrlshRNA SK-N-BE(2)/TGL cells. Same analysis is used as in b.

| miRNA     | Fold-change Stmn Seq2shRNA versus CtrlshRNA | p-value | Target gene(s)       |
|-----------|---------------------------------------------|---------|----------------------|
| Downregulated |                                             |         |                      |
| hsa-miR-3935_st | −2.625                                      | 0.0016  | GABRA4               |
| hsa-miR-1281_st | −1.7572                                     | 0.0056  | MNMAT2               |
| hsa-miR-4668_st | −1.6457                                     | 0.0078  | IBSP, PALM2-2AKAP2   |
| hsa-miR-222_st | −1.624                                      | 0.0317  | NEFH                 |
| hsa-miR-221_st | −1.5368                                     | 0.0057  | INA. NEFH            |
| hsa-miR-620_st | −1.4779                                     | 0.0187  | VGF. C3AR1           |
| hsa-miR-488_st | −1.4515                                     | 0.0099  | GPR158. PLEK1H2     |
| Upregulated  |                                             |         |                      |
| hsa-miR-935_st | 1.4656                                       | 0.0277  | ADAM12. IKZF2. DOCK9. PVRL3. PTPN14 |
| hsa-miR-382_st | 1.5146                                       | 0.0059  | VEGFC. PTPN14       |
| hsa-miR-4656_st | 1.7244                                      | 0.0173  | MARCH3. PTPN14      |
| hsa-miR-132_st | 1.8218                                       | 0.0405  | HUNK. CCDC109B      |
| hsa-miR-4492_st | 2.2542                                       | 0.0308  | STC1                 |

miR-382/PTPN14 expression is deregulated in stathmin-depleted cells. To validate the expression of differentially expressed miRNAs found in the microarrays, qPCR analysis was performed in CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL cells. Of the 12 differentially expressed miRNAs identified, the trends in differential expression between control and stathmin-depleted cells were validated in three miRNAs (miR-488, miR-382 and miR-4656). MiR-488 expression was downregulated, and expression levels of miR-382 and miR-4656 were upregulated by depletion of stathmin using two different siRNAs (Stmn Seq2shRNA and Stmn Seq3shRNA) compared with CtrlshRNA (Supplementary Table 5). PTPN14 was a predicted target for miR-935, miR-382 and miR-4656, the last two miRNAs showing consistent differential expression profiles by both microarray and qPCR (Table 1 and Supplementary Table 5). Moreover, as miR-382 is predicted by both TargetScan and microCosm to target PTPN14 (Supplementary Fig. 4a), we focused on the functional analysis of miR-382/PTPN14. Consistent with the microarray data, PTPN14 mRNA and protein expression were significantly downregulated in Stmn Seq2shRNA and Stmn Seq3shRNA compared with CtrlshRNA cells (Fig. 3a, b). To demonstrate the effect of miR-382 upregulation on PTPN14 expression, we analysed PTPN14 gene and protein expression after transiently transfecting SK-N-BE(2) cells with a miRNA mimic of miR-382. Expression of miR-382 was significantly upregulated with a simultaneous trend to a decrease in PTPN14 gene and protein levels (Supplementary Fig. 4b, c). Conversely, transiently transfecting SK-N-BE(2) cells with a miRNA inhibitor of miR-382 led to decreased expression of miR-382 that was associated with an increase in PTPN14 gene expression (Supplementary Fig. 4d). A trend to downregulate PTPN14 expression was also confirmed in primary tumour samples obtained from stathmin-depleted SK-N-BE(2)/TGL xenografts compared with the CtrlshRNA tumours (Fig. 3c). Collectively, these results demonstrate that expression of miR-382 and its predicted target PTPN14 can be modulated by the levels of stathmin in the neuroblastoma cell line SK-N-BE(2).

Knockdown of PTPN14 expression leads to an increase in neuroblastoma cell migration and invasion. To understand the role of PTPN14 on cellular functions in neuroblastoma cells, we used a gene-silencing approach in two independent neuroblastoma cell lines. The 72-h time point for PTPN14 treatment resulted in maximal knockdown of the protein (Supplementary Fig. 5a). This time point was used in all subsequent experiments for PTPN14. First, we demonstrated that suppression of PTPN14 did not significantly affect SK-N-BE(2)/TGL and SY-SH5Y/TGL cell proliferation (Fig. 4a, b and Supplementary Fig. 5b–e). Using invasion chambers, our results showed that knockdown of PTPN14 increased the migratory ability of SK-N-BE(2)/TGL cells from 43 ± 8% in control cells to 52 ± 6% (p = 0.054) in

Hierarchical clustering of 36 differentially expressed miRNAs in Stmn Seq2shRNA versus CtrlshRNA SK-N-BE(2)/TGL cells. Same analysis is used as in b. To validate the expression of differentially expressed miRNAs found in the microarrays, qPCR analysis was performed in CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL cells. Of the 12 differentially expressed miRNAs identified, the trends in differential expression between control and stathmin-depleted cells were validated in three miRNAs (miR-488, miR-382 and miR-4656). MiR-488 expression was downregulated, and expression levels of miR-382 and miR-4656 were upregulated by depletion of stathmin using two different siRNAs (Stmn Seq2shRNA and Stmn Seq3shRNA) compared with CtrlshRNA (Supplementary Table 5). PTPN14 was a predicted target for miR-935, miR-382 and miR-4656, the last two miRNAs showing consistent differential expression profiles by both microarray and qPCR (Table 1 and Supplementary Table 5). Moreover, as miR-382 is predicted by both TargetScan and microCosm to target PTPN14 (Supplementary Fig. 4a), we focused on the functional analysis of miR-382/PTPN14. Consistent with the microarray data, PTPN14 mRNA and protein expression were significantly downregulated in Stmn Seq2shRNA and Stmn Seq3shRNA compared with CtrlshRNA cells (Fig. 3a, b). To demonstrate the effect of miR-382 upregulation on PTPN14 expression, we analysed PTPN14 gene and protein expression after transiently transfecting SK-N-BE(2) cells with a miRNA mimic of miR-382. Expression of miR-382 was significantly upregulated with a simultaneous trend to a decrease in PTPN14 gene and protein levels (Supplementary Fig. 4b, c). Conversely, transiently transfecting SK-N-BE(2) cells with a miRNA inhibitor of miR-382 led to decreased expression of miR-382 that was associated with an increase in PTPN14 gene expression (Supplementary Fig. 4d). A trend to downregulate PTPN14 expression was also confirmed in primary tumour samples obtained from stathmin-depleted SK-N-BE(2)/TGL xenografts compared with the CtrlshRNA tumours (Fig. 3c). Collectively, these results demonstrate that expression of miR-382 and its predicted target PTPN14 can be modulated by the levels of stathmin in the neuroblastoma cell line SK-N-BE(2).

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PTPN14 expression regulates the YAP/Hippo signalling pathway in a neuroblastoma cell-type-dependent manner. To further elucidate the signalling pathway involving PTPN14 on neuroblastoma cells, we first investigated the impact of PTPN14 siRNA on stathmin expression. We found no significant alteration in stathmin mRNA and protein expression in the PTPN14-depleted SK-N-BE(2)/TGLells (Fig. 5a, b). We then focused on one of the known downstream targets of PTPN14, the Yes-associated protein (YAP or its paralogue Taz), which is a key regulator of the Hippo signalling pathway. In SH-SYSY/TGL cells, only YAP was detected, whereas only its paralogue Taz was expressed in SK-N-BE(2)/TGL cells (Fig. 5c). Our results showed that PTPN14 downregulation did not significantly alter the expression of YAP/Taz compared with the control siRNA-transfected cells (Fig. 5c). The activity of YAP/Taz is mediated by its translocation to the nucleus and subsequent transcription of genes that promoted cell proliferation, migration, invasion and metastasis. We therefore examined the cellular localisation of YAP/Taz in PTPN14-depleted neuroblastoma cells. PTPN14 suppression led to an increase in Taz nuclear translocation, which is significant in PTPN14 Seq4siRNATransfected SK-N-BE(2)/TGL cells compared with the control (Fig. 5d). Analysis of CYR61 mRNA expression, which is one of the target genes of YAP/Taz, showed a significant increased expression in PTPN14-depleted SK-N-BE(2)/TGL cells (Fig. 5e). However, using another neuroblastoma cell line SH-SYSY/TGL, our results showed no significant alteration in YAP nuclear translocation and CYR61 mRNA expression in PTPN14-depleted SH-SYSY/TGL cells compared with the control (Fig. 5f, g). Collectively, these results indicate that PTPN14 and stathmin do not interact in a feed-forward manner and the expression levels of PTPN14 regulate the YAP/Hippo signalling pathway in a neuroblastoma cell line-dependent manner.

**DISCUSSION**

Unravelling the mechanisms of metastasis remains a significant biological challenge to neuroblastoma research and highlights the need for mechanistic studies so that ultimately therapeutic vulnerabilities can be targeted. Here, we provide new insight into stathmin’s molecular mechanism in neuroblastoma metastasis. We identified for the first time a role for stathmin in the modulation of genes and miRNAs in neuroblastoma, leading to the enrichment of genes in pathways associated with functions in cell adhesion, migration and invasion. Focusing on one
predicted target, we unrevealed a new function of the protein PTPN14 as an inhibitor of neuroblastoma cell migration and identified a potential new target for this metastatic disease (Supplementary Fig. 6).

Stathmin is largely recognised for its ability to interact with tubulin and regulate microtubule dynamics. However, this cytoskeletal protein has also been observed to interact with proteins other than tubulin, many of which have been associated with more migratory and invasive cellular phenotypes such as p27Kip1 and STAT3.21,22 Whilst many of the protein functions of stathmin have been reported, its potential role in regulating gene expression and miRNA has been poorly defined. MiRNAs are well known to be a key component driving tumour metastasis in many different cancers including neuroblastoma.12,13,23,24 Here, we identified 12 deregulated miRNAs resulting from the stable knockdown of stathmin expression in the SK-N-BE(2) neuroblastoma cells and validated three of them including miR-382, which has been identified as an important regulator of tumour metastasis.25–28 We found that miR-382 was upregulated in stathmin-depleted cells that we have previously demonstrated to have a reduced metastatic potential.9,10

Our differential and functional gene analysis in stathmin-modulated cells has identified a subset of candidate targets with predicted functions in cell migration and adhesion such as PTPN14,18 PVRL3,29 ADAM1230 and VEGFC.31 These results converged into a global modulation of cell–cell and cell–matrix interactions having essential roles in cell motility and migration, confirming that stathmin expression can modulate aggressive, metastatic neuroblastoma in part by regulation of genes with functions in cell migration and adhesion.

The non-receptor tyrosine phosphatase PTPN14 (also known as Pez, PTPD2 and PTP36) is a developmentally regulated non-receptor protein tyrosine phosphatase (PTP).18 Its role in cancer has recently emerged as a tumour suppressor. Different studies have shown that PTPN14 modulates cell proliferation and invasion.32,33 In addition, a number of its nonfunctional mutations have been

Fig. 3 PTPN14 expression is downregulated in stathmin-depleted SK-N-BE(2)/TGL cells and xenografts. a RT-qPCR analysis was performed on RNA isolated from the CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL cells. The graph shows the quantitative analysis of PTPN14 gene expression normalised to β2-microglobulin. b Representative western blot for PTPN14 protein on whole-cell extracts from the CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL cells. GAPDH was included as a control for equal loading. Graph showing the quantitative analysis of PTPN14 protein expression after normalising to GAPDH. Columns, mean of three independent experiments; bars, SEM. *p < 0.05; **p < 0.01; ****p < 0.0001, statistically significant when comparing the stathmin-depleted cells with the CtrlshRNA. c Histology images of tumours from CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL xenografts with haematoxylin and eosin (H&E), PTPN14 isotype control staining. Graph showing the overall scores for PTPN14 protein expression of CtrlshRNA, Stmn Seq2shRNA and Stmn Seq3shRNA SK-N-BE(2)/TGL primary tumours (n = 4 individual mouse samples/conditions).
reported in breast, pancreas and colorectal tumours. In neuroblastoma, Schramm et al. have found a PTPN14 nonfunctional mutation in neuroblastoma relapse correlating with an aggressive phenotype. However, the biological functions of PTPN14 remain poorly characterised. In this study, we identified PTPN14 as one of the predicted targets for three of the differentially expressed miRNAs (miR-382, miR-935 and miR-4656). Our previous studies have shown that downregulation of stathmin led to a decrease in cell migration and invasion in both in vitro and in vivo neuroblastoma metastasis models. Here, we have found that in stathmin-depleted cells, PTPN14 expression is downregulated, suggesting a compensatory mechanism in stathmin-depleted neuroblastoma cells. To determine its independent role in neuroblastoma, we depleted PTPN14 using siRNA in two independent neuroblastoma cell lines. Our results showed an increase in neuroblastoma cell migration and invasion in the absence of any effect on stathmin levels, indicating that PTPN14 and stathmin did not act in a feedback regulatory loop in the PTPN14-depleted cells. These data suggest that downregulation of PTPN14 expression alone enhances cell migration and invasion in neuroblastoma cells. Given our previous findings in stathmin-depleted neuroblastoma cells that showed a significant reduction in cell migration, invasion and metastatic spread, our current results suggest a complex cross talk between signalling pathways involved in the metastatic process. Further studies to understand the interrelationship between stathmin and PTPN14 expression and the migration/invasion phenotype in neuroblastoma are warranted.

Previous work demonstrated that PTPN14 can suppress YAP/Taz activity by promoting its cytoplasmic localisation. Recently, Cui et al. demonstrated that miR-4516 exerts its oncogenic function by directly targeting PTPN14-mediated regulation of Hippo pathway in glioblastoma. In addition, Schramm et al. showed that expressing the inactive mutant form of PTPN14 caused nuclear translocation of YAP and enhanced clonogenicity of SK-N-SH neuroblastoma cells. Our study
indicates a neuroblastoma cell-type-dependent molecular consequence of PTPN14 inactivity on YAP/Taz signalling, which might rely on the physiological amount of the YAP/Taz in the nucleus. However, given the modest effect on YAP signalling, our findings also suggest that the effects of PTPN14 knockdown on neuroblastoma migration and invasion could be mediated by an alternative pathway. In support of an alternate pathway, a recent study in breast cancer cells showed that PTPN14 can reduce the secretion of a suite of pro-metastatic factors by altering protein trafficking.40

A previous study has reported that PTPN14 regulates cell proliferation in neuroblastoma.37 The lack of effect of PTPN14 on
Fig. 5  PTPN14 expression regulates the YAP/Hippo signalling pathway in a neuroblastoma cell-type-dependent manner. a Relative mRNA expression of stathmin following PTPN14 downregulation in SK-N-BE(2)/TGL cells. Stathmin gene expression normalised to β2-microglobulin. b Representative western blot for equal loading. Graph showing the quantitative analysis of stathmin protein expression following normalisation to GAPDH. c, Representative western blot showing YAP and Taz expression in the PTPN14-depleted SH-SY5Y/TGL and SK-N-BE(2)/TGL cell lines. GAPDH was included as control for equal loading. The human breast adenocarcinoma cell line MCF-7 was included as a positive YAP/Taz staining. d, Representative western blot for Taz protein in SK-N-BE(2)/TGL cells following PTPN14 downregulation. GAPDH was included as a control for equal loading in the cellular lysates (c) and Topo I for the nuclear fraction (N). Graph showing the quantitative analysis of Taz protein translocated in the nucleus. e Relative mRNA expression of CYR61 following PTPN14 downregulation in SK-N-BE(2)/TGL cells. CYR61 gene expression normalised to β2-microglobulin. Columns, mean of at least three independent experiments; bars, SEM. *p < 0.05, **p < 0.01, statistically significant when comparing the PTPN14-depleted cells with the control. f Representative western blot for YAP protein in SH-SY5Y/TGL cells as described in d, g Relative mRNA expression of CYR61 following PTPN14 downregulation in SH-SY5Y/TGL cells as described in e.

CONCLUSIONS
Metastatic neuroblastoma remains a major clinical challenge. Our study provides the first evidence that stathmin can modulate the expression of miRNA and mRNA in neuroblastoma cells, leading to an enrichment of genes involved in migration and invasion signalling pathways. Our data suggest that PTPN14 expression could play a role in the aggressiveness of neuroblastoma and may provide potential new treatment strategies.

ACKNOWLEDGEMENTS
We thank Dr Simon Brayford for critically reviewing the paper.

AUTHOR CONTRIBUTIONS
S.T.P., M.L.G. and M.B.B. wrote the paper, performed the experiments and interpreted the data. A.J.G. participated in analysis of data; G.J.G. and Y.K.G. participated in analysis of data and preparation of the paper. M.K. conceived and supervised the entire study and interpretation of data and editing. All authors read and approved the final paper.

ADDITIONAL INFORMATION
Ethics approval and consent to participate Tumour sections were obtained from control and stathmin-depleted cells xenograft previously obtained in our lab.10 Animal experiments were approved by the Animal Ethics Committee, University of New South Wales (ACEC #13/1168).

Consent to publish N/A.

Data availability The data that support the findings of this study are available on request from the authors.

Competing interests The authors declare no competing interests.

Funding information This work was supported by the Children’s Cancer Institute, which is affiliated with the University of New South Wales (UNSW Australia), and the Sydney Children’s Hospital Network, and by grants from The Kids Cancer Project (MK), National Health and Medical Research (Program Grant APP1091261 and Principal Research Fellowship APP11119152 to MK). MK is also supported by Australian Research Council Centre of Excellence in Convergent Bio-Nano Science and Technology (CE140100036).

Supplementary information is available for this paper at https://doi.org/10.1038/s41416-019-0669-1.

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Cell proliferation in our study may be due to the functionally different methods used for inactivating PTPN14. Our study used RNAi-mediated suppression of PTPN14 expression, whereas Schramm et al. inactivated PTPN14 by mutation.37 Mutated proteins can confer distinct functions on cells that differ from depletion of a protein. To date, no chemical compounds have been developed to target stathmin or PTPN14. Understanding the regulatory mechanisms associated with metastasis has the potential to reveal promising therapeutic strategies against stathmin or PTPN14.
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