

# Analysis of receptor tyrosine kinases (RTKs) and downstream pathways in chordomas<sup>†</sup>

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We have previously demonstrated that chordomas express activated platelet-derived growth factor receptor (PDGFRB) and that treatment with imatinib, which is capable of switching off the activation of various receptor tyrosine kinases (RTKs) including PDGFRB, benefits a number of patients. The aim of this study was to identify the possible presence of other activated RTKs and their downstream signaling effectors. Cryopreserved material from 22 naïve sporadic chordomas was investigated for the presence of activated RTKs and their cognate ligands and downstream signaling effectors by means of human phospho-RTK antibody arrays, Western blotting, and molecular analysis; immunohistochemistry and fluorescence in situ hybridization were used to analyze the corresponding formalin-fixed and paraffin-embedded samples. We detected activated PDGFRB, FLT3, and colony stimulating factor 1 receptor (CSF1R) of the PDGFR family and highly phosphorylated EGFR, HER2/neu, and (to a lesser extent) HER4 of the EGFR family. The detection of PDGFRB/PDGFRB confirmed our previous data. The presence of activated EGFR was paralleled by the finding of high levels of epidermal growth factor (EGF) and transforming growth factor  $\alpha$  (TGF $\alpha$ ) and PDGFRB co-expression and PDGFRB co-immunoprecipitation. Of the downstream effectors, the PI3K/AKT and RAS/MAPK pathways were both activated, thus

leading to the phosphorylation of mammalian target of rapamycin (mTOR) and 4E-BP1 among the regulators involved in translational control. Taken together, our results (i) provide a rationale for tailored treatments targeting upstream activated receptors, including the PDGFR and EGFR families; (ii) support the idea that a combination of upstream antagonists and mTOR inhibitors enhances the control of tumor growth; and (iii) indicate that the 4E-BP1/eIF4E pathway is a major regulator of protein synthesis in chordoma.

**Keywords:** chordomas, mTOR, receptor tyrosine kinases (RTKs), signal transduction.

Chordomas are rare low-grade bone tumors that account for 1%–4% of all malignant bone tumors;<sup>1</sup> they arise from the sacrum in 50%–60% of cases, from the skull base in 20%–35%, from the cervical vertebrae in 10%, and from the thoracolumbar vertebrae in 5%.<sup>2</sup> Their peak incidence is between the fourth and sixth decades of life, although children may also be affected (5%). The M:F ratio is 2–3:1 (more males are affected by sacral chordomas, whereas the frequency of chordomas of the skull base seems to be the same in men and women).<sup>3</sup> Chordomas are generally sporadic, but there have also been familial cases<sup>4–6</sup> and tumors arising in patients with tuberous sclerosis complex (TSC).<sup>7–11</sup>

Untreated disease has an indolent but relentless course. The tumors are locally highly aggressive and may also give rise to distant metastases (10%–30% of cases, or more). On a population basis, the reported 5- and 10-year survival rates are about 70% and 40%, which reflects the naturally slow progress of the disease; however, after the development of metastases, median survival is about 1 year.<sup>3</sup>

Surgery remains the best standard treatment for both localized and metastatic disease; radiotherapy can

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control tumor growth, particularly cobalt or heavy particle radiation, but the high doses required lead to significant toxicity, and standard cytotoxic chemotherapy is inefficacious.<sup>3</sup> Targeted molecular therapy recently been found to be promising as imatinib has induced an anti-tumor response in a number of patients when used alone<sup>12</sup> and when combined with cisplatin in patients showing secondary progression on imatinib monotherapy.<sup>13</sup> Furthermore, combined treatment with imatinib and sirolimus has proved to be effective in patients with imatinib-resistant chordomas.<sup>14</sup>

We have previously demonstrated that chordomas express activated platelet-derived growth factor receptor (PDGFRB) and that this activation is due to the occurrence of an autocrine/paracrine loop;<sup>15</sup> it has also been reported that a subset of chordomas express EGFR and c-MET.<sup>16</sup> The presence of activated receptor tyrosine kinases (RTKs) leads to the activation of secondary transducers belonging to the MAPK or PI3K/AKT pathways, which together activate mTOR;<sup>17</sup> downstream of mTOR, S6K/S6, and 4E-BP1 harmonize protein synthesis, and promote cell growth and proliferation.

Recent insights into the regulation of translation control have shown that, in addition to the p70 ribosomal S6 kinase (p70S6K)-mediated PI3K/mTOR pathway, S6 may also be regulated by the p90 ribosomal S6 kinase (p90RSK)-mediated RAS/MAPK cascade.<sup>18</sup> Furthermore, 4E-BP1 (a repressor of eIF4E) may be inactivated as a result of hyper-phosphorylation by the PI3K/mTOR or RAS/MAPK cascade through a still unidentified mechanism (which abrogates the association between 4E-BP1 and eIF4E),<sup>19</sup> or as a result of transcription silencing<sup>20,21</sup> (Fig. 1). More importantly, S6 may act as an oncogene or tumor suppressor gene.<sup>22,23</sup> Finally, it is likely that the specific functions of the 2 regulators of the translational machinery are closely related to cell type or tumor histotype.<sup>24</sup>

In this study, human phospho-RTK antibody arrays (RTK cards) were used to explore the upstream RTK profile of 7 naïve sporadic chordomas, and detected activated PDGFRB, FLT3, and colony stimulating factor 1 receptor (CSF1R) of the PDGFR family, as well as highly phosphorylated EGFR, HER2/neu, and (to a lesser extent) HER4 of the EGFR family. Taken together, this subsequently validated upstream analysis indicates that both the PDGFR and EGFR families may be activated in chordomas. This upstream analysis was extended to 22 sporadic cases, and downstream effectors were investigated. We found activated PI3K/AKT and RAS/MAPK cascades lead to the phosphorylation of mTOR. The absence of gene deregulation upstream of mTOR in the presence of PDGFRB and EGFR ligands supports the hypothesis that the activation is ligand dependent and that combined RTK and mTOR/MAPK inhibitors<sup>25</sup> may lead to better disease control. Our findings also indicate that translational control downstream of mTOR is preferentially driven by the 4E-BP1/eIF4E pathway.

## Materials and Methods

### *Clinical Data and Materials*

We analyzed 21 surgical specimens and 1 wide biopsy sample taken from 22 surgically treated patients with a confirmed diagnosis of conventional sporadic chordoma: 16 males and 6 females with a median age of 61 years (range 36–77). Fourteen were primary tumors and 8 were recurrences (Patients 7, 10–12, 14, 19, 20, and 22); 18 were located in the sacrum and 4 in the thoraco-lumbar vertebrae. Freshly frozen material and formalin-fixed paraffin-embedded (FFPE) blocks were available for all but 1 of the patients (Patient 14), for whom there was only frozen material.

All of the analyses involved representative areas of tissue sections carefully selected under microscope control in order to avoid normal tissue or necrosis. The biochemical and molecular analyses were performed using representative cryopreserved tissue and checking 1 hematoxylin-and-eosin-stained frozen section; the immunohistochemical (IHC) and fluorescence in situ hybridization (FISH) analyses were performed using the corresponding FFPE samples.

Written informed consent was obtained from all of the patients.

### *Analyses of Frozen Material*

**Biochemical Analysis.**—*RTK Phosphorylation Array.* The expression of phosphorylated growth factor RTKs was detected using the Proteome Profiler Array Kit (ARY001, R&D Systems) as laid down in the manufacturer's protocol, and 2 mg of protein lysate per array.

**Total Protein Extraction, Immunoprecipitation and Western Blotting:** The proteins were extracted from tissue samples stored at  $-80^{\circ}\text{C}$  as previously described.<sup>26</sup> For the immunoprecipitation (IP) analysis, 1 mg each of protein lysates was precipitated by means of incubation with Protein A Sepharose (Amersham Biosciences) and specific antibodies: 20  $\mu\text{l}$  of rabbit polyclonal anti-PDGFRB (sc-432, Santa Cruz Biotechnology), 10  $\mu\text{l}$  of rabbit polyclonal anti-EGFR (sc-120, Santa Cruz Biotechnology), and 10  $\mu\text{l}$  of mouse monoclonal anti-Her2/Neu (Ab-3, Calbiochem; MGR2 kindly provided by Dr. Tagliabue, Fondazione IRCCS Istituto Nazionale dei Tumori, Milan, Italy).

Immunoblotting was performed using the anti-phosphotyrosine mouse monoclonal antibody (05–321, Upstate) diluted 1:3000 to reveal the absence or presence of activated/phosphorylated receptor. The filters were stripped and then incubated with the specific antibody in order to detect the degree of receptor expression: anti-PDGFRB (sc-432, Santa Cruz Biotechnology), anti-EGFR (sc-03, Santa Cruz Biotechnology), and anti-HER2/neu (Ab-3, Calbiochem). The filter of IP anti-PDGFRB was stripped again and incubated with anti-PI3-kinase p85 $\alpha$  (sc-1637, Santa Cruz Biotechnology); the filters of IP anti-EGFR were incubated with anti-PDGFRB and/or anti-HER2/neu in order to detect the co-immunoprecipitated proteins.

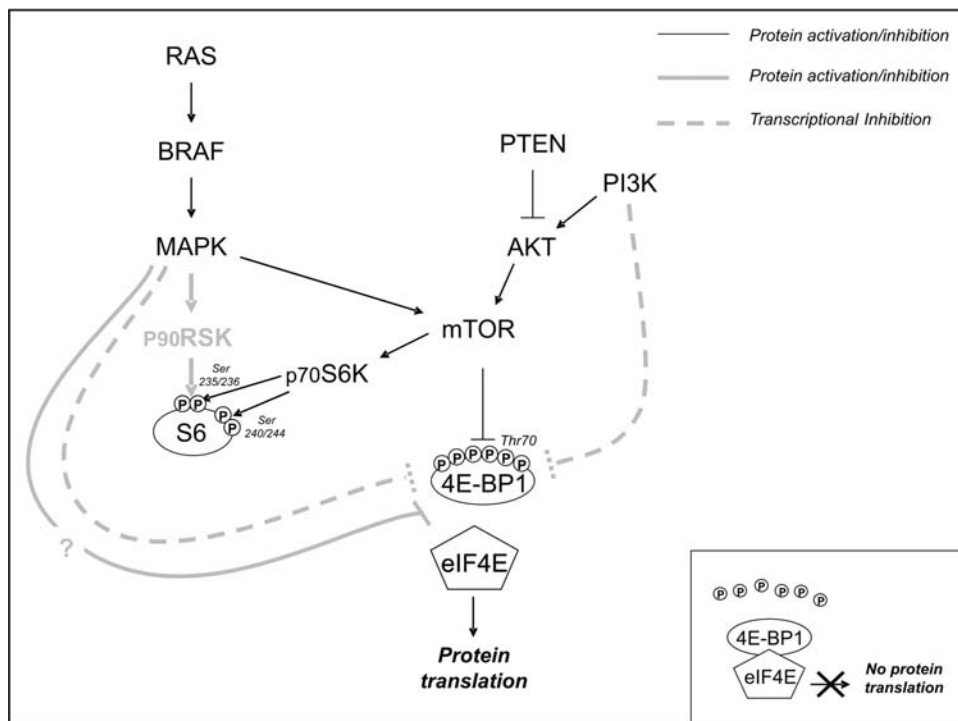


Fig. 1. Usual regulation of RTK downstream effectors. The grey lines indicate new pathways related to protein activation/inhibition by means of phosphorylation (thick continuous lines) or transcriptional silencing (dotted lines). In addition to being phosphorylated by p70S6K/mTOR, S6 may be activated by p90RSK. The same is true of 4E-BP1 which, in addition to mTOR, may be phosphorylated by an unknown factor (light-grey thick continuous lines); it may also be regulated by the transcriptional silencing induced by MAPK and/or PI3K (light-grey thick dotted lines). Insert: Non-phosphorylated 4E-BP1 forming a complex with eIF4E that blocks protein translation.

For the EGFR IP/WB experiments, the protein bands detected on filters were densitometrically analyzed on X-ray film images using Quantity One Software (Bio-Rad). The densitometrically determined levels of phosphorylated EGFR were calculated as ratios of phosphorylated and expressed EGFR.

The expression and activation of the downstream targets were detected by means of direct Western blotting (WB) after loading 20 µg of total protein/lane using the antibodies anti-P-AKT (#4058S Ser473, Cell Signaling Technology), anti-AKT (#9272, Cell Signaling Technology), anti-P-MAPK (#4376S Thr202/Tyr204, Cell Signaling Technology), anti-MAPK (#9102 p44-42, Cell Signaling Technology), anti-P-mTOR (#2971S Ser2448, Cell Signaling Technology), anti-mTOR (#2972, Cell Signaling Technology), anti-P-S6 (#2215 Ser240/244 and #2211 Ser 235/236, Cell Signaling Technology), anti-P-4E-BP1 (#9455 Thr70, Cell Signaling Technology), anti-4E-BP1 (#9644 Cell Signaling Technology), anti-PTEN (#9559, Cell Signaling Technology), and anti-actin (A2066, Sigma). The negative control for PTEN WB was a cell line without PTEN protein expression (3962 M, kindly provided by Dr. Rodolfo, Fondazione IRCCS Istituto Nazionale dei Tumori).<sup>27</sup>

**RNA Extraction and Real-Time PCR.**—Total RNA was extracted from fresh frozen tissue and reverse transcribed as previously described.<sup>28</sup>

**RTK Ligand Detection.** The presence or absence of EGFR ligand mRNA [EGF and transforming growth factor α (TGFα)] was verified by means of real-time PCR using specific EGF (Hs00153181\_M1) and TGFα probes (Hs00608187\_M1) (Applied Biosystems); Table 1 shows the experimental conditions. The mRNA was not quantified.

**PTEN and S6.** Real-time PCR of *PTEN* and *S6* was performed on respectively, 4 and 6, representative samples using specific *PTEN* (Hs02621230\_S1) and *S6* probes (Hs01058685\_g1), and the experimental conditions shown in Table 1.

**DNA Extraction and Sequencing.**—DNA was digested by proteinase-K and underwent standard phenol extraction. Mutation analyses were made of exons 18–21 of *EGFR*, exons 9 and 20 of *PI3KCA*, exons 5–9 of *PTEN*, exons 1 and 2 of *KRAS*, and exons 11 and 15 of *BRAF*. Table 1 shows the primers and experimental conditions used for PCR amplification. After purification using the ExoSAP-IT Kit (USB Amersham), the DNA of the PCR products was automatically sequenced (ABI prism 377; Applied Biosystems).

**Analyses of FFPE Material**

**RNA Extraction and RT-PCR.**—Total RNA from microdissected samples was isolated by means of the

**Table 1.** Experimental condition**Real-time PCR**

Ligands		Probes		Conditions		
EGF	Hs00153181_M1 (Applied Biosystem)	50°C 2'–95°C 10'–95°C 15', 60°C 1' × 40 cycles				
TGF $\alpha$	Hs00608187_M1 (Applied Biosystem)					
PTEN	Hs02621230_S1 (Applied Biosystem)					
S6	Hs01058685_g1 (Applied Biosystem)					
Molecular analysis						
Gene	Exon	Sense	Antisense	Annealing temperature (°C)		
<i>EGFR</i>	18	ctgaggtgaccctgtctc	tacagcttgcaaggactctg	60 × 40 cycles		
	19	agatcactgggcagcatgtg	cagctgccagacatgagaa			
	20	ctctggccaccatgcgaa	ccaggcaactctgctatcc			
	21	tcttccatgatgatctgt	cctgggtgcaggaaaatgct			
PI3K	9	gggaaaaatagacaagaaagc	ctgagatcagccaaattcagtt	58 × 35 cycles		
	20	ctcaatgatgctggctctg	tggaatccagatgagctttc			
PTEN	5	tgcaacatttctaaagttactact	gaggaaaggaaaaacatcaaaa	55 × 40 cycles		
	6	ttttcaattggcttctctttt	tgttccaatacatggaaggatg			
	7	cagttaaaggcatttctctgtg	ttttgatatttctccaatgaa			
	8	tgcatattcattcttttctttc	aagtcaacaacccccacaaa			
K-Ras	1	tggtggagtattgatagtg	catgaaaatggtcagaga	56 × 35 cycles		
	2	ggtgcactgtaataatccagac	tgatttagtattattatggc			
B-Raf	11	tcctctcaggcataaggtaa	cgaacactgaatatttctttgat	55 × 40 cycles		
	15	tcataatgcttctgatagga	ggcaaaaattaatcagtga			
Immunohistochemical analysis						
Antibody	Clone	Company	Dilution	Antigen retrieval	Positive control	
PDGFRB	sc-339	Santa Cruz Biotechnology	1:100	6' at 95°C mM citrate buffer, pH 6	Human breast carcinoma	
EGFR	K1492	Dako	Procedures performed in accordance with manufacturer's protocol			
Her2	Polyclonal	Dako	1:2000	6' at 95°C, pH 6	Human breast carcinoma	

TGF $\alpha$ , transforming growth factor  $\alpha$ .



TRIzol method (Invitrogen), and reverse transcribed using Superscript reverse transcriptase (Invitrogen) and both oligo dT and random hexamers.

**Ligand Detection:** The presence of PDGFB mRNA was verified by means of RT-PCR as previously described.<sup>28</sup>

**Immunohistochemistry.**—The antibodies and experimental conditions are shown in Table 1. The score for each marker is listed below.

**PDGFRB.** Cytoplasmic immunoreactivity was evaluated and scored on the basis of the percentage of immunoreactive cells (not less than 70%). PDGFRB staining intensity was evaluated using vessel wall immunoreactivity as an in-built control.

**EGFR.** The score was applied according to the manufacturer's instructions, combining the scores relating to staining intensity and the percentage of immunopositive cells.<sup>29</sup> The resulting low, intermediate, or high/strong immunodecoration are reported in Table 2. In the cases scored as negative, none of the cells were immunodecorated.

**HER2/neu.** The same parameters considered for PDGFRB were applied for this receptor. As for EGFR, in the cases scored as negative, none of the cells were positive.

For all these markers, the inter-observer correlation for the assessment of the staining was concordant (100%).

**Fluorescence in Situ Hybridization.**—Fluorescence in situ hybridization analyses were used to investigate the gene status of *EGFR*, *HER2/neu*, *PTEN*, and *S6* as previously described.<sup>26,30,31</sup> In particular, the BAC clones RP11-126H20 and RP11-624N8 were used as probes for *S6*, and Spectrum Green(SG)-labeled CEP9 (Vysis) was used as the control probe. The dual-color FISH mixture was prepared using Spectrum Orange (SO)-labeled *S6*+ SG-labeled CEP9. The corresponding gene status was evaluated as previously described.<sup>32</sup> To simplify the FISH/IHC correlations, the high EGFR and HER2/neu polysomy/amplification group was classified as FISH positive (FISH+), and the remaining groups (disomy, trisomy, or low polysomy) as FISH negative (FISH-), whereas the *PTEN* and *S6* gene status patterns and loss/gain percentages were detailed in each case.

## Results

The profiles of the 8 recurrent tumors were almost identical to those of the 14 primary tumors. All of the data are summarized in Table 2.

### Upstream Targets

**RTK Array Assays.**—The RTK array experiments used the 7 chordomas for which adequate total protein

extract was available (Patients 1, 4–6, 8, 12, and 13). RTK activation was evaluated by comparing the intensity of the labeled spots relating to each tumoral protein extract with that of those relating to the positive and negative controls on the array membrane.

All 7 patients showed activated PDGFRB (although the level of activation was low in case Patient 13) and, to a lesser extent, FLT3 and CSF1R of the PDGFR family. EGFR was highly activated in 4 patients (Patients 1, 4, 12, and 13), activated in 2 (Patients 5 and 6), and negative in 1 (Patient 8). HER2/neu was detected in 5 cases (Patients 1, 4, 5, 12, and 13), and HER4 in 3 (Patients 4, 5, and 12). Of the “non-druggable” RTKs, Axl and Dtk were activated in all of the patients. The negative controls were always undetectable. Figure 2A shows 4 representative cases (Patients 4, 5, 12, and 13). Overall, the profiles of the 42 RTKs evaluated on the array membrane indicated significant activation of members of the PDGFR family (PDGFRB, FLT3, and CSF1R) and the EGFR families (EGFR, HER2/neu, and HER4).

### Confirmatory Biochemical Analysis of PDGFRB, EGFR, and HER2/Neu.

—All of the case material was analyzed by means of IP/WB in order to confirm the RTK array data. PDGFRB was activated in all but 1 case, in line with our previous data,<sup>15</sup> whereas EGFR was activated in 17 and not activated in 5 (Patients 1 and 8–11); Fig. 2B shows 12 representative cases. The results obtained using the 2 methods perfectly overlapped, except in Patient 1. The EGFR WB results were quantified by densitometry in order to define the activation level of the receptor more precisely (Fig. 2C).

Total protein extracts from 6 cases expressing activated EGFR (Patients 2, 4, 12, 13, 15, and 18) were immunoprecipitated with anti-EGFR and then incubated with anti-PDGFRB: PDGFRB co-immunoprecipitated with EGFR in all cases (Fig. 2D).

Immunoprecipitation and Western blotting analysis of HER2/neu using antibody directed against the NH<sub>2</sub> terminal showed that all of the cases were negative, which is consistent with the unrecognizable structural conformation of the receptor's extracellular domain. However, an antibody specific for the C-terminal applied to the 14 cases for which material was still available revealed HER2/neu expression/phosphorylation in 4 cases (Patients 4, 5, 15, and 18) and HER2/neu expression alone in 2 (Patients 17 and 20). The IP/WB and array data matched in 4 cases (Patients 4–6 and 8) and did not match in 2 (Patients 12 and 13); insufficient material prevented a comparison in 1 case (Patient 1).

The stripped membranes with EGFR immunoprecipitated proteins were re-incubated with a HER2/neu antibody, which revealed the presence of HER2/neu in 9/22 cases, thus suggesting that the 2 members of the family may be involved in heterodimers. The result was confirmed by further incubation of the stripped filter with the EGFR/PDGFRB co-IP experiment (Fig. 2D, upper panel) and with anti-HER2/neu antibody (Fig. 2D, lower panel).

**Table 2.** Summary results

Cases	Upstream marker and method													
	PDGFRB			PDGFB	EGFR				EGF	TGF $\alpha$	Her2/neu			
	RTK array	IP/WB	IHC	RT-PCR	RTK array	IP/WB	IHC	FISH	Real-time PCR	Real-time PCR	RTK array	IP/WB	IHC	FISH
1	+	+P	+	+	+	-	-	-	+	+	+	-	-	
2		+P	+	+		+P	-	-	+	+		-	-	
3		+P	+	+		+P	+M	-	+	+		-	-	
4	+	+P	+	+	+	+P	+S	+	+	+	+	+P	-	+
5	+	+P	+	+	+	+P	-	NE	+	+	+	+P	-	NE
6	+	+P	+	+	+	+P	-	+	+	+	-	-	-	+
7		+P	+	+		+P	-	+	+	+		-	-	+
8	+	+P	+	+	-	-	-	-	+	+	-	-	-	-
9		+P	+	+		-	-	-	+	+		-	-	-
10		+P	+	+		-	-	-	+	+		-	-	-
11		+P	+	+		-	NE	-	+	+		-	-	-
12	+	+P	+	+	+	+P	-	NE	+	+	+	-	-	NE
13	+	+P	+	+	+	+P	+S	-	+	+	+	-	-	-
14		+P	+	+		+P	-	-	+	+		-	-	-
15		+P	+	+		+P	-	-	+	+		+P	-	-
16		-	+	+		+W	-	-	+	+		-	-	-
17		+P	+	+		+W	-	-	+	+		+P	-	-
18		+P	+	+		+P	+L	-	+	+		-	-	-
19		+P	+	+		+P	+S	-	+	+		-	-	+
20		+P	+	+		+P	+M	-	+	+		+	-	-
21		+P	+	+		+P	+S	NE	+	+		-	NE	NE
22		+P	+	+		+P	-	-	+	+		-	-	-

*Continued*

**Table 2.** *Continued*

Cases	Downstream marker and method												
	P85 IP/WB	AKT WB	PTEN WB	FISH	Real-time	ERK 1/2 WB	mTOR WB	S6 WB	PS6 Thr240/244 WB	PS6 Thr235/236 WB	S6 FISH	Real-time	4EBP1 WB
1	+	+P	+	M(80)	+	+P	+P	+	-	±P	D	+	+P
2	+	+P	+	D	+	+P	+P	+	±P	±P	D		+P
3	+	+P	+	M(70)	+	+P	+P	+	±P	±P	D	+	+P
4	+	+P	+	HP		+P	+P	+	+P	+P	D	+	+P
5	+	+P	+	NE		+P	+P	+	±P	+P	NE		+P
6	+	+P	+	HP(10)		+P	+P	-	-	-	D	+	+P
7	+	+P	+	M(50)		+P	+P	±	-	±P	D		+
8	+	+P	+	D	+	+P	-	-	-	-	LP	+	+P
9	+	-	-	D		+P	±P	-	-	-	D		+
10	+	+P	+	M(50)		+P	+P	+	±P	-	D	+	+P
11	+	+P	+	D		+P	+P	-	-	-			+
12	+	+P	+	M(50)		+P	+P	+	±P	+P			+P
13	+	+P	+	D		+P	+P	±	-	±P	D		+P
14	+	+P	+			+P	+P	+	±P				+
15	+	+P	+	D		+P	+P	+	±P		M		+P
16	-	+P	+	D		+P	+P	-	-	-	D		+
17	+	+P	+	M(60)		+P	+P	+	±P		NE		+
18	+	+P	+	D		+P	+P	-	-	-	D		+P
19	+	+P	+	D		+P	+P	-	-	-	D		±
20	+	+P	+	M(90)		+P	+P	+	±P		D		+P
21	+	+P	+	NE		+P	+P	-	-	-	NE		-
22	+	+P	+	D		+P	+P	+	±P				+

The empty spaces correspond to not performed analyses. Abbreviations: IP, immunoprecipitation; WB, Western blotting; IHC, immunohistochemistry; FISH, fluorescence in situ hybridization; NE, not evaluable; P, phosphorilated; W, weak; D, disomic; HP, high polisomy (percentage); M, monosomic (percentage). For EGFR IHC: S, strong; M, moderate; L, low.

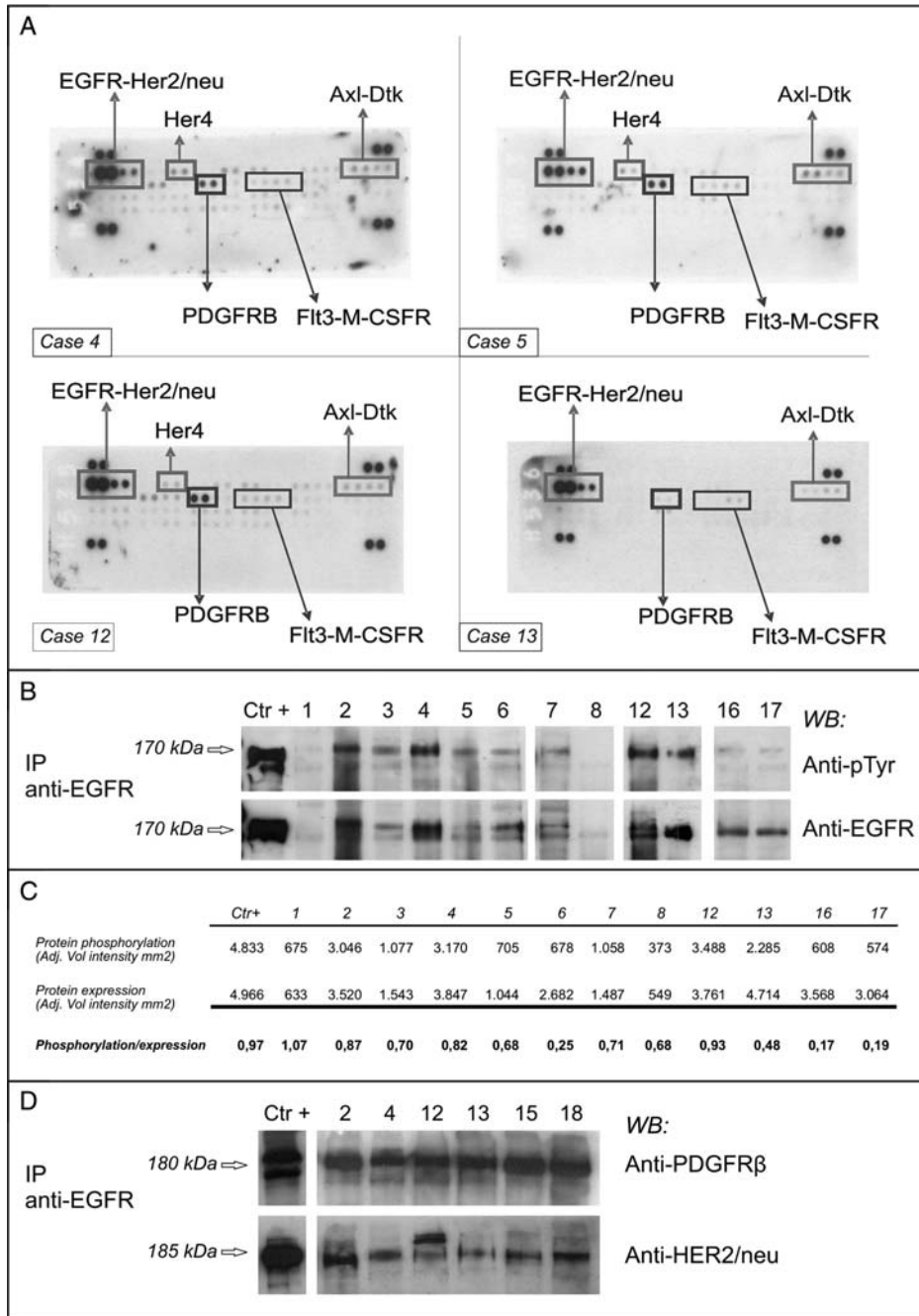


Fig. 2. Expression and activation of RTKs. The sample numbers correspond to the cases listed in Table 2. (A) RTK arrays. Equal amounts of total protein extracts from 4 representative cases were incubated with the arrays. The rectangles indicate the presence of activated receptors. (B) EGFR IP/Western blot analysis of 12 representative cases. To confirm the presence of activated EGFR, proteins were immunoprecipitated with a specific antibody and blotted onto a membrane. The anti-pTyr panel identified the phosphorylated EGFR; the anti-EGFR panel indicates the expression of the corresponding receptor. The A431 cell line (American Type Culture Collection, Manassas, VA) was used as the positive control. (C) Densitometric analysis of levels of EGFR activation. The bands representing phosphorylated EGFR and total EGFR levels were quantified, and the ratios between phosphorylated and expressed EGFR were calculated, the ratio relating to the A431 cell line (positive control) was 0.97. The low level of phosphorylation detected in samples #16 and #17 may have been due to methodological problems (non-optimal cryopreservation). (D) Co-IP of EGFR/PDGFRB and EGFR/HER-2/neu. Total protein extracts from 6 EGFR-positive cases were immunoprecipitated with anti-EGFR antibody and run on a gel. The filter was incubated with anti-PDGFRB antibody, then stripped and re-incubated with anti-HER2/neu antibody. Extracts from the 2N5A cell line (immunoprecipitated with anti-PDGFRB) and SKBR3 cell line (immunoprecipitated with anti-HER2/neu) were respectively used as positive controls for PDGFRB and HER2/neu protein. The 2N5A cell line (derived from the NIH3T3 cell line and expressing the COL1-PDGFB fusion that characterized dermatofibrosarcoma protuberans) was kindly provided by Dr. Greco, Fondazione IRCCS Istituto Nazionale dei Tumori; the SKBR3 came from the American Type Culture Collection.



**Analysis of the Mechanisms Sustaining PDGFRB and EGFR Activation.**—We have previously demonstrated the absence of PDGFRB-activating mutations in chordomas;<sup>15</sup> in this study, we analyzed the EGFR gene to exclude a mutation-based activation mechanism. EGFR was wild type in all of the samples.

Real-time PCR showed that all of the cases highly expressed both EGFR ligands (EGF and TGF $\alpha$ ) and as expected,<sup>15</sup> PDGFB mRNA was expressed in all of the analyzed cases (Table 2). We thus concluded that, most likely, the activation of these 2 receptors is sustained by an autocrine/paracrine loop.

### Downstream Effectors

We biochemically and/or molecularly investigated the PI3K/AKT (the p85 and p110 subunits of PI3K, AKT, and PTEN) and MAPK pathways (RAS, BRAF, and Erk1/2), and then analyzed the mTOR kinase and its S6 and 4E-BP1 downstream effectors. The analysis of PTEN and S6 was complemented by the FISH analysis of FFPE material.

All of the data are summarized in Table 2.

**PI3K/AKT Pathway.**—As a first confirmatory step of receptor activation, we verified the binding of the p85 subunit of PI3K to activated PDGFRB: co-IP was detected in all of the samples. The p110 subunit gene (*PI3KCA*) was investigated by looking for activating mutations, and the hot spots corresponding to exons 9 and 20 were sequenced; no mutations were detected in any of the cases.

Western blotting showed that AKT was activated to different extents in all of the analyzed samples (Fig. 3A upper panel), except Patient 9.

PTEN phosphatase reverses the effects of p110-PI3K. We sequenced the mutation hot spot (from exon 5 to exon 9) but did not detect any alterations.

As gene copy number is 1 of the mechanisms responsible for PTEN destabilization, we used FISH to analyze matched FFPE samples and found alterations in 9 (47%) of the 19 successfully analyzed cases: chromosome 10 monosomy in 7 cases (1 of which is shown in Fig. 3B) and a high degree of polysomy in 2 cases. However, WB evaluating the overall expression of PTEN protein indicated no differences in expression levels (Fig. 3C; the results were validated using a cell line null for PTEN expression). Given the discrepancy between the FISH analysis and protein expression, we further analyzed *PTEN* mRNA expression in 2 disomic (Patients 2 and 8) and 2 monosomic cases (Patients 1 and 3); the biochemical results were confirmed by real-time PCR as no differences were found between the monosomic and disomic cases (Fig. 4D).

**MAPK Pathway.**—Analysis of the *KRAS* and *BRAF* mutation hot spots revealed no activating mutations.

Direct WB showed that both p44 and p42 (Erk1/2) were much more highly expressed and phosphorylated than any other of the analyzed proteins (Fig. 3A lower panel).

**mTOR, S6, and 4E-BP1.**—mTOR is activated by both pathways and, as both cascades are activated in chordomas, we expected to find an activated kinase in all 22 samples. WB revealed a phosphorylated protein in all but 2 cases (Patient 9 was only weakly positive; Patient 8 was negative) (Fig. 4A).

The activation status of S6 was investigated using Ser235–236 and Ser240–244 phospho-antibodies. Ser240–244 residues are preferentially phosphorylated in response to the mTOR/PI3K pathway, and S6 was found to be expressed and activated to a low/very low extent in 11 cases, not expressed in 8, and expressed but not phosphorylated in 3 (Fig. 4B). The same trend (a low level of activation) was observed in the 13 cases that were analyzed using the Ser235–236 antibody, which also detects the residues phosphorylated in response to the mTOR-independent RAS/MAPK pathway.

As the low level of protein expression/activation despite mTOR activation suggested that S6 plays an oncosuppressing role, we used FISH (on FFPE material) and real-time PCR to ascertain its gene status and mRNA expression. The FISH results showed that all but 1 of the analyzed cases had a disomic or low polysomic pattern (Patient 15 was monosomic, with positive protein expression). However, it is possible that our probe does not detect the very small 3979 bp deletion of the S6 gene,<sup>33</sup> and that the loss in Patient 15 (which also carried monosomy of the p16 gene closely mapping on the same chromosome; data not shown) was attributable to a longer deletion encompassing both genes, although a second probe gave the same results (data not shown). Nevertheless, real-time PCR showed the same level of S6 transcript in the samples with (Patients 1, 3, 4, and 10) and without S6 protein expression (Patients 6 and 8).

To investigate 4E-BP1, we used the Thr70 phospho-antibody to highlight protein hyper-phosphorylation, which, by abrogating the association between 4E-BP1 and eIF4E, releases translational repression. 4E-BP1 was hyper-phosphorylated in 13 cases, not expressed in 1, and expressed but not phosphorylated in 8 (Fig. 4C), which means that eIF4E was released in 14 cases, thus favoring protein synthesis, cell proliferation, and tumor progression. In 4 cases (Patients 9, 11, 16, and 19), 4E-BP1 was expressed but not phosphorylated (and thus eIF4E was repressed), and the fact that S6 was null supports the hypothesis that malignant progression may be sustained by the absence of S6, which may act as an oncosuppressing gene as it does in chondrosarcomas.<sup>23</sup> The level of S6 activation was low in the remaining 4 cases in which eIF4E was repressed (Patients 7, 14, 17, and 22). In Patient 8, in which 4E-BP1 was hyper-phosphorylated in the absence of mTOR activation and Ser235–236/240–244 S6 phosphorylation, it is possible to assume activation through the RAS/MAPK cascade mediated by a still unknown factor,<sup>19</sup> although many oncogenic events may lead to 4E-BP1 phosphorylation.<sup>34</sup>

Taken together, our findings downstream of mTOR suggest that both PI3K/AKT and RAS/MAPK signaling are involved in regulating protein transduction and that the 4E-BP1 pathway drives protein synthesis, whereas

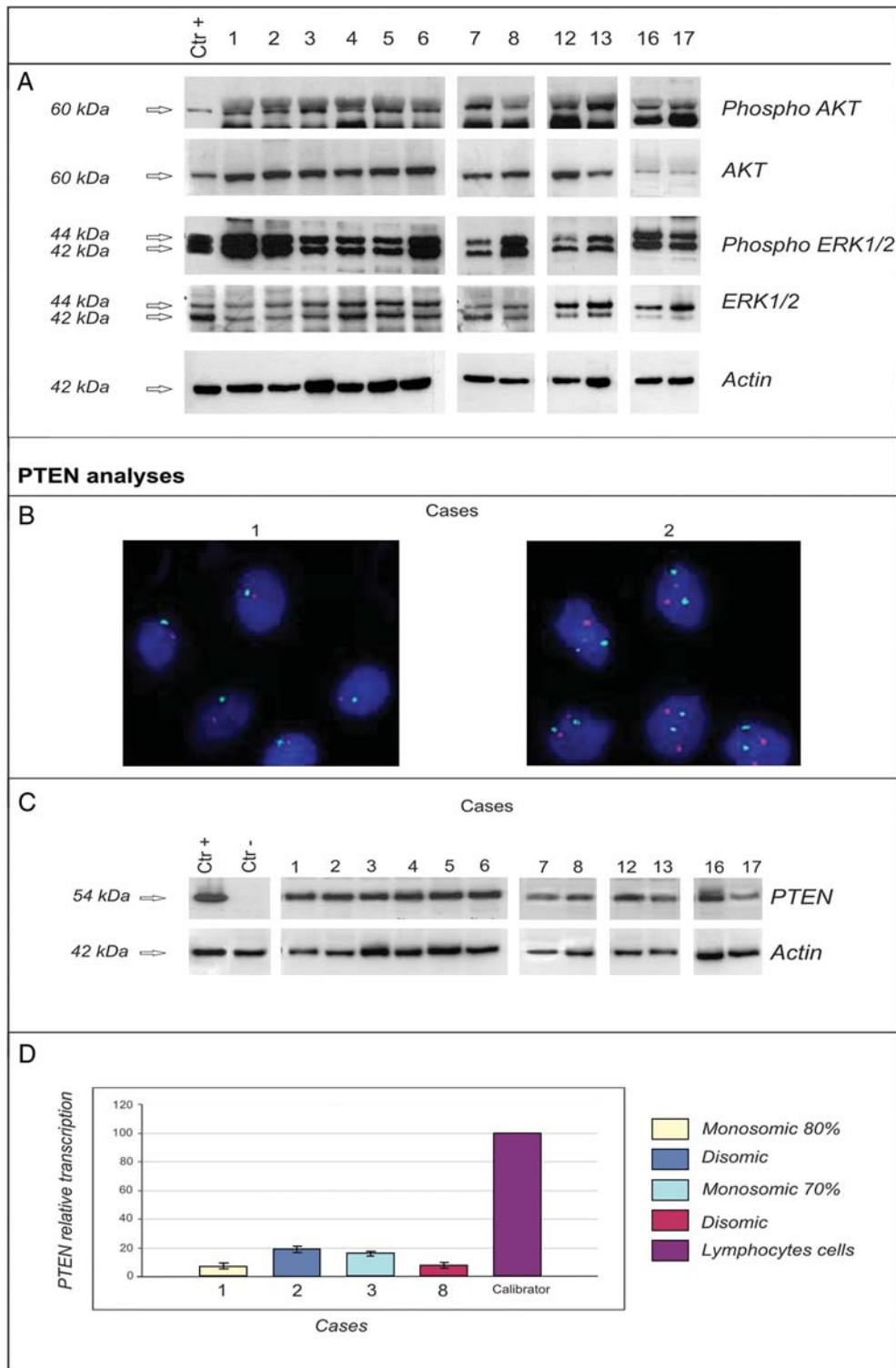


Fig. 3. Expression of downstream signaling markers. (A) Western blots showing the presence of phosphorylated/activated and total AKT and ERK1/2. PTEN analyses: (B) FISH. The chromosome 10 probe is labeled with Spectrum Green and the *PTEN* locus-specific probe with Spectrum Orange. Case 1: monosomic *PTEN* in 80% of the cells. Case 2: disomic *PTEN*. (C) Western blotting. *PTEN* expression in 12 representative monosomic or disomic cases: monosomic cases were Patients 1, 3, 7, 12, and 17. The different electrophoretic mobility observed in Patient 16 may have been due to protein hyper-phosphorylation, which cannot be confirmed by the antibody used because it does not discriminate between phosphorylated and non-phosphorylated *PTEN*. The A431 and 3962 M cell lines were, respectively, used as positive and negative controls, and anti-actin antibody was used to normalize the results. (D) Real-time PCR in the disomic and monosomic cases. Lymphocyte mRNA was used for calibration purposes.

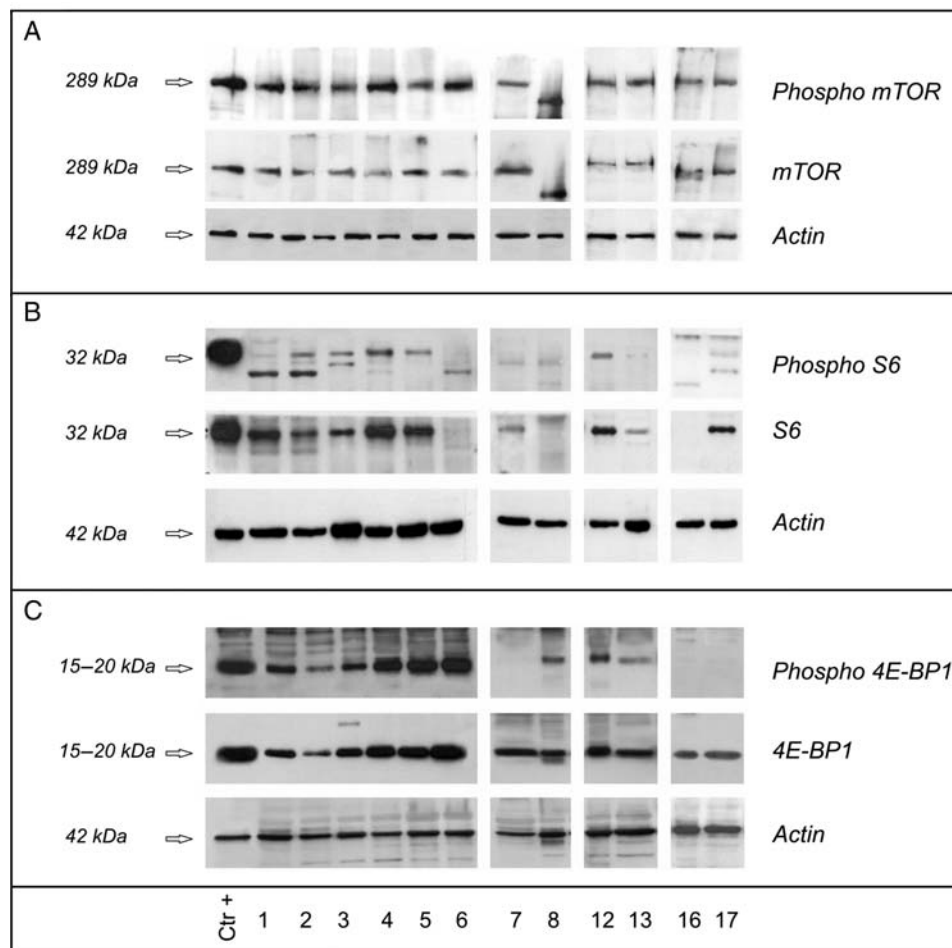


Fig. 4. mTOR expression and its downstream effectors. mTOR (panel A), S6 (panel B), and 4E-BP1 (panel C) in the same 12 representative cases analyzed and reported in Figure 3, panel A. Twenty micrograms of total protein extract was loaded for each sample. The A431 cell line was used as the positive control and anti-actin antibody was used to normalize the results.

the hypothesized oncosuppressing role of S6 remains undetermined and will require further experiments.

**Methods of Selecting Patients for Possible Targeted Treatments.**—Our previous positive experience with PDGFRB immunophenotyping in chordoma,<sup>15</sup> and the recent favorable results obtained in non-small cell lung cancer using gene and protein profiling,<sup>35,36</sup> prompted us to find a screening method capable of identifying predictive markers for TK inhibitors that can be easily applied to FFPE samples (frozen material is not frequently available). To this end, we investigated the sensitivity of IHC and FISH analyses (alone or combined), taking the biochemical results as the gold standard.

**IHC.** In all the cases (100%), more than 80% of the cells showed PDGFRB cytoplasmatic immunoreactivity, with the intensity of the staining being stronger than that of the vessel wall used as the control.

EGFR was expressed in 7/20 cases (4 classified as strong, 2 as moderate, and 1 as low) showing a decoration mainly related to the cell membrane. In the

remaining cases, scored as negative, none of the tumoral cells were positive for EGFR.

All cases were negative for HER2/neu as all tumor cells were negative.

**FISH.** As amplification is one of the mechanisms responsible for receptor activation in the case of EGFR family genes, we used FISH to analyze the EGFR and HER2/neu genes. Of the 18 evaluable cases, the EGFR gene was disomic in 11 and low polysomic in 4 (Patients 10 and 17–19) (FISH–) and high polysomic in 3 (Patients 4, 6, and 7) (FISH+).

The HER2/neu gene was disomic in 10 and low polysomic in 4 (Patients 10, 11, 17, and 18) (FISH–), and was high polysomic in 3 (Patients 4, 7, and 19) or amplified in some areas (Patient 6) (FISH+).

All 7 cases with deregulated EGFR also showed deregulated HER2/neu; they all also had the same deregulation pattern except for Patients 19 and 6.

**Correlations Between IP/WB, IHC, and FISH.**—EGFR. We compared the IP/WB results with those of IHC and FISH in 17 evaluable patients. The IP/WB and

IHC data correlated in 9 cases: in particular, all of the IP/WB-negative cases were confirmed by IHC, whereas only 5 of the 13 IP/WB-positive cases (38%) were also IHC positive, thus indicating that IHC is less sensitive than IP/WB but does not generate false positives.

The combination of IHC and FISH increased the number of “targetable” patients by recovering 2 FISH+ cases from the 8 IHC-negative and IP/WB-positive patients (7/13 cases; 54%).

*HER2/neu*. This receptor was identified by IP/WB in 6/14 evaluable cases (43%), all of which were IHC negative. The lack of any correlation between a gain in gene copy number and *HER2/neu* expression suggests that IP/WB positivity may be due to heterodimer formation.

## Discussion

The aim of this study was to use biochemical analyses (RTK array cards that can provide an overview of phosphorylated RTK in small amounts of total protein extracts and IP/WB) of cryopreserved material to demonstrate and validate the presence of a reproducible panel of activated receptors from mainly the 2 RTK families of PDGFR and EGFR. We have previously shown that PDGFRB is activated by means of a ligand-dependent mechanism;<sup>15</sup> we now confirm the PDGFRB data and show that the pathogenesis of chordoma also involves EGFR (Table II, IP/WB column, Fig. 2B and C).

The co-expression and co-immunoprecipitation of PDGFRB and EGFR has already been described at the preclinical level, and it has been demonstrated that PDGFRB stimulates the activation of PDGFRB and subsequently stimulates EGFR unidirectionally.<sup>37,38</sup> Our results demonstrate the co-IP of EGFR and PDGFRB in surgical specimens and suggest heterodimer formation. If these findings are confirmed, they would explain the existence of imatinib-insensitive chordomas which, as they are also addicted for EGFR, can be expected to escape imatinib treatment. It is interesting to note in this regard that a response to cetuximab (an EGFR-specific monoclonal antibody) has been reported in 2 chordoma patients.<sup>39,40</sup>

In addition to the interaction between the 2 families, our co-IP data also indicate the possible existence of heterodimerization between individual members of the EGFR family at least in some cases, as supported by the finding that the receptor was only bound by a specific C-terminal antibody, and not by an N-terminal antibody. This suggests that the use of bi-specific inhibitors such as lapatinib, which blocks both EGFR and *HER2/neu*,<sup>41–43</sup> may offer more clinical benefit than mono-specific treatments.

Finally, the phospho-RTK-array approach has shown that other members of the PDGFR family are activated, including CSF1R and FLT3, which have been reported to be inhibited by imatinib.<sup>44,45</sup>

After the upstream RTK analysis, we biochemically and/or molecularly and cytogenetically investigated

some downstream effectors, and the results indicated that both the PI3K/AKT and MAPK pathways were activated in the absence of activating mutations. The co-IP experiments showed that the p85 subunit of PI3K directly binds the activated/dimerized PDGFRB receptor and, in line with this, AKT was phosphorylated (Fig. 3A). Molecular and FISH analyses of the *PTEN* oncosuppressor gene showed the absence of inactivating mutations and the presence of *PTEN* alterations (mainly chromosome 10 monosomy) in 7 of 19 cases, although the latter do not seem to affect protein function because direct WB did not reveal any difference in the expression of the phosphatase between the monosomic and disomic cases (Fig. 3B and C). The WB results were further confirmed by real-time PCR. We also detected highly expressed and phosphorylated ERK1 and ERK2 (Fig. 3A), which is consistent with activation of the EGFR family, in the presence of wild-type KRAS and BRAF genes (but these 2 markers were not biochemically analyzed). Finally, in line with previously published data indicating that both AKT and ERK1/2 can activate mTOR, the protein was phosphorylated in all but 1 case (Fig. 4A), which suggests that it may be a further “druggable” target.

Taken together, our results show that mTOR activation is due to the growth factor-mediated activation of PI3K and MAPK, with no contribution from PTEN. This means that our findings only partially overlap those of a recent WB study of the growth-factor deprived, chordoma-derived U-CHI cell line and IHC investigations of sporadic sacral chordomas samples, on the basis of which the authors suggested that partial or complete PTEN deficiency may be responsible for mTOR signaling activation.<sup>46</sup>

Although needing further confirmation, our findings concerning the 2 regulators of the translational machinery downstream of mTOR raise important speculative and practical issues. Unlike that in alveolar sarcomas,<sup>47</sup> protein synthesis in chordomas seems to be harmonized by 4E-BP1 rather than S6, and this reinforces the idea that the translational apparatus may differently regulate protein production (and thus the cell cycle and tumor progression) in different histotypes.<sup>24</sup> It is currently believed that the 4E-BP1/eIF4E pathway plays a key role in tumorigenesis and the 4E-BP1 is a “funnel effector” that is relevant to prognosis and treatment response.<sup>34</sup> Our finding that the 2 effectors have selective and histotype-restricted functional roles raise doubts about the use of the phospho-S6 antibody as a biomarker for the activation of the PI3K/mTOR pathway in everyday practice, and indicates a need to extend the analysis to the phospho-4E-BP1 antibody.

Finally, the low level of S6 activation suggests that S6 may act as an oncosuppressor gene, even if the experimental proof here reported do not indicate any gene copy loss or any decrease in mRNA levels in the protein-negative cases.

As previous IHC investigations have shown that PDGFRB predicts receptor activation in 100% of cases,<sup>15</sup> we analyzed matched FFPE samples for EGFR and *HER2/neu* by means of IHC and interphase FISH



(taking the biochemical results as the gold standard) in an attempt to acquire insights into identifying patients who may benefit from targeted EGFR and HER2/neu treatments. The results indicated that after combining IHC and FISH (2 techniques that are commonly used in the case of FFPE material), EGFR sensitivity increases to 54% without any false positives and so, despite the constant HER2/neu null immunophenotype (given the co-expression of EGFR and HER2/neu), this may be sufficient for patient selection. As HER2/neu is involved in EGFR dimer formation, its null immunophenotype is expected and has previously been reported in chordomas. Cumulatively, our results are in agreement with previously reported IHC data<sup>48</sup> for PDGFRB and Her2/neu, while a lower percentage of positivity was observed for EGFR. This discrepancy, however, could be due to the different experimental conditions applied (different EGFR antibody).

In conclusion, our data indicate new avenues for combining chordoma treatments<sup>49</sup> by targeting upstream and downstream effectors. In addition to the PDGFR family, which is inhibited by imatinib, the presence of activated receptors of the EGFR family may provide a

rationale for treating these patients with small molecules that are effective on both EGFR and HER2/neu. Moreover, the strong activation of downstream signaling supports the idea that a combination of upstream TK antagonists and mTOR and MAPK<sup>25</sup> inhibitors could lead to the better control of tumor growth by avoiding the so-called negative feedback loop,<sup>17</sup> lapatinib resistance,<sup>50</sup> and the PI3K-dependent feedback loop.<sup>25</sup> However, further studies are needed to clarify the multifaceted regulation of downstream mTOR effectors in different histotypes.

*Conflict of interest statement.* None declared.

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