

## MEN1 Gene Mutation and Reduced Expression Are Associated with Poor Prognosis in Pulmonary Carcinoids

Dorian R.A. Swarts<sup>1\*</sup>, Aldo Scarpa<sup>2†</sup>, Vincenzo Corbo<sup>2</sup>, Wim Van Criekinge<sup>3</sup>, Manon van Engeland<sup>4</sup>, Gaia Gatti<sup>5</sup>, Mieke E.R. Henfling<sup>1</sup>, Mauro Papotti<sup>5</sup>, Aurel Perren<sup>6</sup>, Frans C.S. Ramaekers<sup>1</sup>, Ernst-Jan M. Speel<sup>4†</sup>, Marco Volante<sup>5</sup><sup>[GRAPHIC]</sup>

Departments of <sup>1</sup>Molecular Cell Biology and <sup>5</sup>Pathology, GROW – School for Oncology & Developmental Biology, Maastricht University Medical Centre, Maastricht, The Netherlands; <sup>2</sup>ARC-NET Research Centre and Department of Pathology and Diagnostics, University and Hospital Trust of Verona, Verona, Italy; <sup>3</sup>Laboratory of Bioinformatics and Computational Genomics, Ghent University, Belgium; <sup>4</sup>Department of Oncology, Division of Pathology, University of Turin at San Luigi Hospital, Orbassano, Torino, Italy; <sup>6</sup>Department of Pathology, University of Bern, Switzerland

**Context:** *MEN1* gene alterations have been implicated in lung carcinoids, but their effect on gene expression and disease outcome are unknown.

**Objective:** To analyse *MEN1* gene and expression anomalies in lung neuroendocrine neoplasms (NENs) and their correlations with clinicopathologic data and disease outcome.

**Design:** We examined 74 lung NENs including 58 carcinoids and 16 high-grade neuroendocrine carcinomas (HGNECs) for *MEN1* mutations (n=70) and allelic losses (n=69), promoter hypermethylation (n=65), and mRNA (n=74) expression. Results were correlated with disease outcome.

**Results:** *MEN1* mutations were found in 7/55 (13%) carcinoids and in 1 HGNEC, mostly associated with loss of the second allele. *MEN1* decreased expression levels correlated with the presence of mutations ( $P=0.0060$ ) and was also lower in HGNECs than carcinoids ( $P=0.0024$ ). *MEN1* methylation was not associated with mRNA expression levels. Patients with carcinoids harbouring *MEN1* mutation and loss had shorter overall survival ( $P=0.039$  and  $P=0.035$ , respectively), and low *MEN1* mRNA levels correlated with distant metastasis ( $P=0.00010$ ) and shorter survival ( $P=0.0071$ ). In multivariate analysis, stage and *MEN1* allelic loss were independent predictors of prognosis.

**Conclusion:** Thirteen percent of pulmonary carcinoids harbour *MEN1* mutation, associated with reduced mRNA expression and poor prognosis. Also in mutation-negative tumours, low *MEN1* gene expression correlates with an adverse disease outcome. Hypermethylation was excluded as the underlying mechanism.

**P**ulmonary neuroendocrine neoplasms (NENs) include well differentiated carcinoids, classified into typical (TCs) and atypical (ACs) subtypes, with a relatively favorable prognosis (1, 2), and aggressive high-grade neuroendocrine carcinomas (HGNECs), ie, large cell neuroendocrine carcinoma (LCNEC) and small cell lung cancer (SCLC).

Lung carcinoids occur in 5% of patients with multiple endocrine neoplasia (MEN) type 1 syndrome with *MEN1* germline mutations (3). *MEN1* mutations have been reported in approximately 16% of sporadic lung carcinoids

Lung carcinoids occur in 5% of patients with multiple endocrine neoplasia (MEN) type 1 syndrome with *MEN1* germline mutations (3). *MEN1* mutations have been reported in approximately 16% of sporadic lung carcinoids

ISSN Print 0021-972X ISSN Online 1945-7197

Printed in U.S.A.

Copyright © 2013 by The Endocrine Society

Received July 8, 2013. Accepted November 12, 2013.

Abbreviations: AC, atypical carcinoid; FISH, fluorescent *in situ* hybridisation; HGNEC, high-grade neuroendocrine carcinoma; LCNEC, large cell neuroendocrine carcinoma; MBD, methyl-CpG-binding domain; MSP, methylation-specific MSP; NEN, neuroendocrine neoplasia; NGS, next-generation sequencing; qRT-PCR, quantitative real-time PCR; SCLC, small cell lung cancer; TC, typical carcinoid.

(4–6), but are exceptional in HGNECs (7, 8). Allelic losses of the *MEN1* locus are detected more frequently (~36%) than gene mutations (5), suggesting that alternative mechanisms might lead to *MEN1* gene inactivation. So far, the association of *MEN1* gene alterations with clinical follow-up has not been reported in lung NENs.

Here we report a comprehensive analysis of *MEN1* alterations, methylation status and transcriptional expression in 74 lung NENs. Results are compared with tumor characteristics and clinical outcome.

## Materials and Methods (detailed in Supporting Information)

### Tissue samples

Seventy-four sporadic lung NENs (34 TCs, 23 ACs, 1 unclassified carcinoid, 9 LCNECs, 6 SCLCs and 1 mixed LCNEC-SCLC), including 71 with follow-up (median 52 months) were studied. Institutional review board approval for the study was obtained by each contributing center following local ethical requirements. Nine neuroendocrine cell lines were included for *MEN1* mRNA expression analysis.

### *MEN1* mutational analysis

The entire coding region and splice junctions of *MEN1* were analyzed on tumor and matched normal DNA by Sanger sequencing, as described (9) (Supplementary Table 1A).

### *MEN1* deletion analysis

Fluorescent in situ hybridization on frozen sections was performed using the *MEN1* locus BAC-probe CTD-222019, and centromere probes for chromosomes 1, 11, 3 and 7 as described (10).

### Quantitative real-time PCR (QRT-PCR)

QRT-PCR was performed using mRNA on *MEN1*, menin interactors *JUND* and *MLL*, and menin targets *HOXC8*, *P18*, *P21*, *P27*, using primers listed in Supplementary Table 1B. Expression levels were normalized to *ACTB* and *CYP19A1* housekeeping genes as described (11).

### Methylation-Specific PCR (MSP) and next-generation sequencing (NGS)

*MEN1* promoter methylation analysis was performed on all cases by nested MSP (Supplementary Table 1C) and by NGS on Methyl-CpG-binding Domain (MBD)-immunoprecipitated DNA of 10 primary carcinoids and carcinoid cell lines NCI-H720 and NCI-H727 (focusing on the *MEN1* gene).

### Statistical analysis

Correlations between *MEN1* data and follow-up were determined using IBM SPSS Statistics (v20.0.0; IBM) setting the *P*-values cut-off for significance at  $\leq 0.05$ . Differences in mean *MEN1* mRNA levels were determined by Student's *t* test, and the optimal cut-off point was determined using area under the ROC-curve analysis. Associations between clinical and molecular pa-

rameters were calculated using logistic and linear regression, Fisher's exact or  $\chi^2$  tests, as appropriate. Associations between relative expression levels of *MEN1* and other genes were calculated using Pearson's correlation.

Survival curves were created by the Kaplan-Meier method and compared using the log-rank test. The prognostic value of age at diagnosis, histopathology, sex, smoking status, tumor diameter and *MEN1* mutation, deletion and gene expression were tested in univariate and multivariate settings using Cox-regression. Only variables with a *P*-value  $< 0.10$  in univariate analyses were included in multivariate analyses.

## Results

### *MEN1* gene mutations and deletions are more frequent in ACs

*MEN1* mutations were searched in 70 tumors, including 7 previously published cases (5). Eight cases (11%) showed nonsynonymous somatic alterations: 2/32 TCs (6%), 5/23 ACs (22%) and 1/7 LCNECs (14%). Mutations included three small deletions, three single nucleotide substitutions and two insertions (Supplementary Tables 2 and 3). patient 35 showed the same *MEN1* mutations in the primary tumor and two distant metastases removed eight and ten years later.

Ploidy status and *MEN1* deletion were analyzed by FISH in 69 tumors (Table 1B and Figure 1A-C). Most carcinoids were disomic whereas HGNECs were more frequently polysomic. Among carcinoids, 8/57 (14%, 1 TC and 7 ACs) showed *MEN1* allelic loss. In HGNECs, *MEN1* was deleted in 3/12 cases (25%). *MEN1* deletion was more frequent in ACs than in TCs ( $P = .0060$ ), and was significantly associated with presence of mutations in both the total group of NENs ( $P = .0022$ ) and in carcinoids ( $P = .0063$ ) (Supplementary Table 3).

### *MEN1* gene expression is heterogeneous in lung NENs

Quantitative RT-PCR performed on all 74 NENs showed that relative *MEN1* expression levels in lung NENs ranged from 0.031 to 0.85, generally lower than exocrine pancreas (mean 0.58; Figure 1D) but higher than normal lung (mean 0.078). A gradual decrease in mean *MEN1* expression values was apparent from TCs to ACs and HGNECs (Figure 1D-E). Similarly, *MEN1* expression levels in SCLC NCI-H69 and LCNEC NCI-H460 cell lines were lower than in lung carcinoid NCI-H720 and NCI-H727 cells (Figure 1D). *MEN1* gene expression levels were inversely correlated to the presence of mutations and deletions (Supplementary Table 3) in the complete group of lung NENs ( $P \leq .0060$ ) and carcinoids ( $P \leq .025$ ).

**Table 1.** Univariate and multivariate analysis of clinical data and *MEN1* parameters correlated with outcome of patients with pulmonary neuroendocrine neoplasms (NENs)

	A) Lung NENs		B) Carcinoids	
	Univariate <i>P</i> -value	Multivariate* <i>P</i> -value	Univariate <i>P</i> -value	Multivariate* <i>P</i> -value
Age at diagnosis	<b>0.0003</b>	NS	<b>0.0021</b>	NS
Diameter	0.086	NI	NS	NI
Histopathology	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>0.011</b>	NS
Sex	<b>0.049</b>	NS	NS	NI
Smoking status	NS	NI	NS	NI
Stage	<b>0.0002</b>	NS	<b>0.0013</b>	<b>0.0009</b>
<i>MEN1</i> deletion	<b>0.013</b>	NS	<b>0.047</b>	<b>0.013</b>
<i>MEN1</i> mutation	0.060	NI	0.050	NI
<i>MEN1</i> gene expression	<b>0.0068</b>	<b>0.017</b>	<b>0.035</b>	NS

\*Only parameters displaying a *P*-value <0.05 are included in multivariate analysis and in the final model all *P*-values needed to be <0.05. Abbreviations used: NEN: neuroendocrine neoplasm; NI, not included in multivariate analysis; NS, not significant

### No correlation between *MEN1* methylation and gene expression

As low *MEN1* expression levels were present in tumors with neither *MEN1* mutation nor deletion, we analyzed *MEN1* promoter methylation by MSP. No methylation-specific product was seen in 65 primary NENs, cell lines and 15 normal lungs from noncancer patients. NGS in 10 primary carcinoids and lung carcinoid cell lines NCI-H720 and NCI-H727 showed some methylation signals in *MEN1* promoter region and gene body, but not at the locus used for MSP analysis (Supplementary Figure 1). No methylation signals were found in CpG-islands. There were no differences in methylation between samples with high and low *MEN1* mRNA levels.

### Deregulation of *MEN1* is associated with metastatic disease and poor prognosis

*MEN1* gene mutations and deletions and decreased gene expression were associated with the presence of distant metastasis in both the total group of NENs and the carcinoids subgroup, but were not related to sex, age and smoking status (Supplementary Table 4). In particular, *MEN1* mRNA levels were significantly lower in patients with distant metastasis and/or patients who died from their tumor, both in the entire group (0.162 vs. 0.318, *P* = .00080) and within the carcinoid subgroup (Figure 1E).

Univariate survival analysis showed in both the complete group of NENs and the carcinoid subgroup that age, histopathology, stage, presence of *MEN1* deletions and decreased gene expression were significantly associated with higher risk of death (Table 1). At multivariate survival analysis, histopathology and decreased *MEN1* gene expression in lung NENs overall, and stage and *MEN1* deletion in the carcinoid subgroup retained statistical significance. Kaplan-Meier survival analysis confirmed that the 15-year overall survival of lung carcinoids was significantly shorter in patients with *MEN1* mutation (Figure

1F), decreased mRNA levels (Figure 1G) and deletion (*P* = .035).

### Menin interactors and target genes

Within the complete set of NENs, *MEN1* gene expression was correlated with that of its interactors *JUND* (*P* = .016) and *MLL* (*P* = .026) but not of its targets *HOXC8*, *P18*, *P21* and *P27*. Significant associations between *MEN1* deletion and low *HOXC8* expression (*P* = .010) and low *P18* gene expression (*P* = .017) were also found. *MEN1* mutational status and menin protein expression were not associated with mRNA levels of interactors and downstream targets.

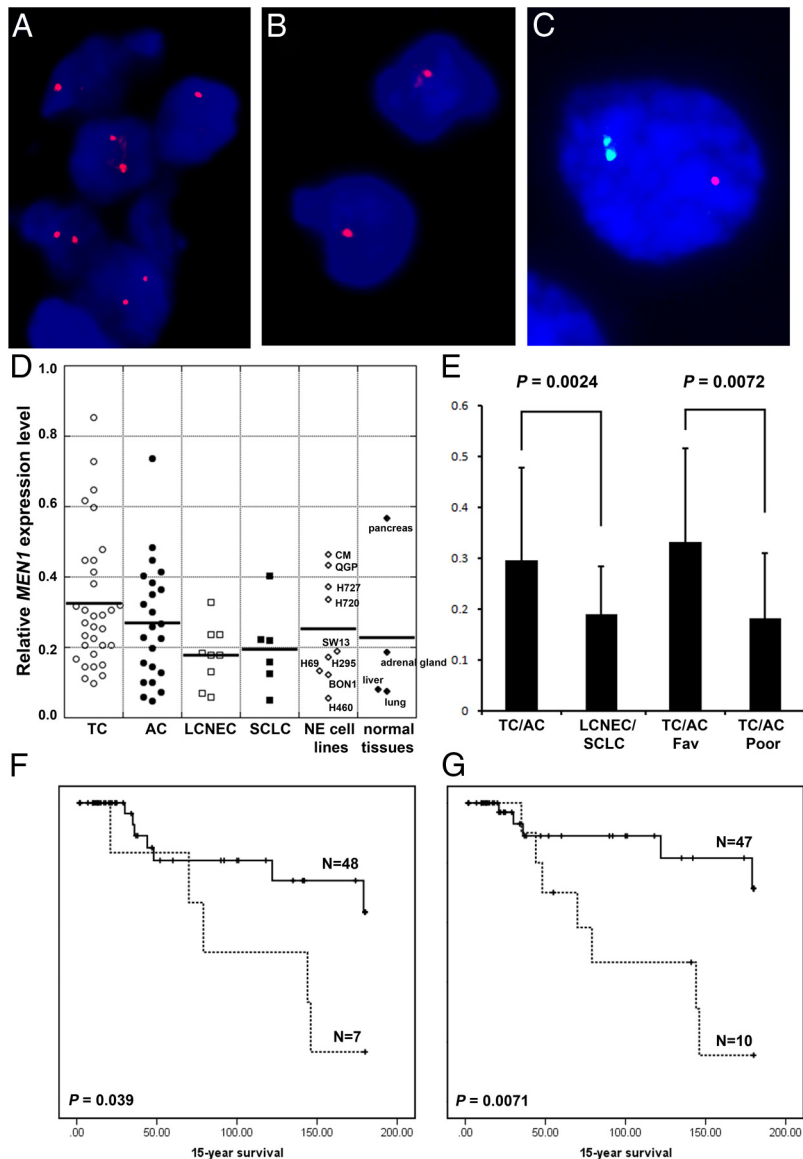
In the complete group of NENs, as well as within carcinoids, lower levels of *P18* expression (*P* = .0074 and *P* = .0011, respectively) were associated with lower 15-year survival.

### Discussion

Our study on pulmonary NENs shows that the presence of *MEN1* mutation and/or a lower *MEN1* gene expression is associated with distant metastasis and lower 15-year survival, in both the entire cohort of NENs and lung carcinoids considered separately.

*MEN1* mutations were found in 13% (7/55) of carcinoids, in line with previous reports (4, 5), with most mutations occurring in ACs. All mutations were of the loss of function type, and most mutated cases also displayed deletion of the second *MEN1* locus. Only 1/15 HGNECs was mutated, confirming that *MEN1* mutations do not play a major role in HGNECs (7, 8).

The large majority of neoplasms with wild-type *MEN1* gene had high mRNA levels. However, not all cases with low mRNA levels harbored mutations or deletions, indicating that other mechanisms might unbalance *MEN1* ex-



**Figure 1.** *MEN1* gene alterations and expression in lung neuroendocrine neoplasms, and association with survival. A-C: Fluorescence in situ hybridization for the *MEN1* locus (in red), showing a case with a *MEN1* mutation but retention of two copies of the *MEN1* gene (A), and two cases with *MEN1* mutation and associated loss of the *MEN1* locus (B and C), the latter in the presence of two copies of chromosome 7 (centromeric probe of chromosome 7 in green). D: Scatterplots showing the relative mRNA expression values of *MEN1* for typical carcinoids (TC), atypical carcinoids (AC), large cell neuroendocrine carcinomas (LCNEC), small cell lung cancers (SCLC), neuroendocrine (NE) cancer cell lines and normal tissues, compared to the geometric mean of the expression levels of the housekeeping genes *ACTB* and *CYPA*. The horizontal bars represent the mean expression values. E: Differences between the mean values  $\pm$  SD of *MEN1* gene expression levels between the total group of carcinoids (TC/AC) and NE carcinomas (LCNEC/SCLC) (left; 95% confidence interval (CI) [CI]  $-0.175/-0.0400$ ) and of carcinoid patients (TC/AC) with a favorable (Fav) and a poor disease outcome (right; 95% CI  $0.0423/0.257$ ). F-G: Kaplan-Meier analyses depicting 15-year overall survival in the group of carcinoids according to the presence of *MEN1* gene mutations (F, dotted line) or *MEN1* gene expression levels below the cut-off point of 0.136 (G, dotted line).

pression in these tumors. No hypermethylation was seen at CpG-islands within the *MEN1* promoter at MSP-analysis, while methylated regions outside the CpG-island were identified by MBD-sequencing. However, such methylation was not associated with reduced *MEN1*

mRNA levels, suggesting that *MEN1* hypermethylation unlikely underlies *MEN1* downregulation.

*MEN1* gene alterations and reduced expression were significantly associated with tumor stage, presence of distant metastases, and patient survival. At multivariate analysis, decreased *MEN1* gene expression was an independent predictor of disease outcome in both the complete group of NENs and in the group of carcinoids alone. This finding seems specific for lung NENs since it is not observed in pancreatic NENs (10, 14).

The mechanisms underlying the role of *MEN1* gene loss of function and the acquisition of clinical aggressiveness in lung carcinoids deserves further investigation. We identified correlations between *MEN1* and *JUND/MLL* mRNA levels, but not with menin targets *P18*, *P21* and *P27*, suggesting that lung-specific mechanisms might be active, such as the E-cadherin/ $\beta$ -catenin pathway which interacts with menin and is frequently deregulated in lung NENs (7, 15, 16).

In conclusion, our comprehensive overview of *MEN1* molecular alterations in lung NENs demonstrates a significant association between *MEN1* gene deregulation and clinical aggressiveness in lung carcinoids and postulates different mechanisms of gene regulation in well and poorly differentiated NENs.

## Acknowledgments

We would like to thank the following people who have provided us with tumor material: Jos Broers, Department of Molecular Cell Biology, Maastricht University, The Netherlands; Thomas Rudolph, Department of Pathology, University of Bern, Switzerland; Matthijs van Oosterhout and Kees Seldenrijk, Department of Pathology, Sint Antonius Hospital, Nieuwegein; Loes van Velthuysen, Division of Pathology, Netherlands Cancer Institute, Amsterdam, The Netherlands. In addition we like

to thank Kim Wouters and Jürgen Veeck, Department of Pathology, Maastricht University Medical Centre, for help with the MSP and the Biobix genome browser, respectively. We also thank Annick Haesevoets, Department of Molecular Cell Biology and Pathology, Maastricht University Medical Centre, for technical assistance, Robert-Jan van Suylen, Department of Pathology, Jeroen Bosch Hospital, 's-Hertogenbosch, and Erik Thunnissen, Department of Pathology, Free University Medical Centre, Amsterdam, for performing reclassification of carcinoid tumors, and Anne-Marie Dingemans, Department of Pulmunology, Maastricht University Medical Centre, Maastricht, The Netherlands, for acquiring clinical follow-up data.

Address all correspondence and requests for reprints to: †Prof. Aldo Scarpa, MD PhD, ARC-NET Research Centre and Department of Pathology and Diagnostics, University of Verona - Policlinico G.B. Rossi, Piazzale L.A. Scuro 10, 37134 Verona, Italy, Phone: +39-458124043, Fax: +39-458127432, Email: aldo.scarpa@univr.it.

\*Current affiliation: Department of Biology, University of York, York, United Kingdom

<sup>[GRAPHIC]</sup>These senior authors contributed equally to the manuscript

**Conflict of interest statement:** W. Van Criekinge is employed as CSO in MdxHealth., E.-J. M. Speel has honoraria from Speakers Bureau of Pfizer and Lilly and is a consultant/advisory board member of Pfizer.

This work was supported by grants from the Italian Association for Cancer Research (AIRC) (grant #IG 13567 to MV; grant n. 12182 to AS), the Italian Cancer Genome Project (FIRB RBAP10AHJB to AS) and a grant from the Jan Dekkerstichting & Ludgardine Bouwmanstichting (nr. 2010 010 to DRAS and EJMS).

## References

1. Beasley MB, Thunnissen FB, Hasleton PhS et al. Carcinoid tumour. In: Travis WD, Brambilla E, Muller-Harmelink HK, Harris CC, editors. Pathology and genetics of tumours of the lung, pleura, thymus and heart. Lyon: IARC Press, 2004: 59–62.
2. Granberg D, Eriksson B, Wilander E, Grimfjard P, Fjallskog ML, Oberg K, Skogseid B. Experience in treatment of metastatic pulmonary carcinoid tumors. *Ann Oncol.* 2001;12:1383–1391.
3. Sachithanandan N, Harle RA, Burgess JR. Bronchopulmonary carcinoid in multiple endocrine neoplasia type 1. *Cancer.* 2005;103: 509–515.
4. Debelenko LV, Brambilla E, Agarwal SK, Swalwell JI, Kester MB, Lubensky IA, Zhuang Z, Guru SC, Manickam P, Olufemi SE, Chandrasekharappa SC, Crabtree JS, Kim YS, Heppner C, Burns AL, Spiegel AM, Marx SJ, Liotta LA, Collins FS, Travis WD, Emmert-Buck MR. Identification of MEN1 gene mutations in sporadic carcinoid tumors of the lung. *Hum Mol Genet.* 1997;6:2285–2290.
5. Görtz B, Roth J, Krahenmann A, de Krijger RR, Muletta-Feurer S, Rutimann K, Saremaslani P, Speel EJ, Heitz PU, Komminoth P. Mutations and allelic deletions of the MEN1 gene are associated with a subset of sporadic endocrine pancreatic and neuroendocrine tumors and not restricted to foregut neoplasms. *Am J Pathol.* 1999; 154:429–436.
6. Veschi S, Lattanzio R, Aceto GM, Curia MC, Magnasco S, Angelucci D, Cama A, Piantelli M, Battista P. Alterations of MEN1 and E-cadherin/beta-catenin complex in sporadic pulmonary carcinoids. *Int J Oncol.* 2012.
7. Debelenko LV, Swalwell JI, Kelley MJ, Brambilla E, Manickam P, Baibakov G, Agarwal SK, Spiegel AM, Marx SJ, Chandrasekharappa SC, Collins FS, Travis WD, Emmert-Buck MR. MEN1 gene mutation analysis of high-grade neuroendocrine lung carcinoma. *Genes Chromosomes Cancer.* 2000;28:58–65.
8. Haruki N, Yatabe Y, Travis WD, Nomoto S, Osada H, Nakamura S, Nakao A, Fujii Y, Takahashi T. Characterization of high-grade neuroendocrine tumors of the lung in relation to menin mutations. *Jpn J Cancer Res.* 2000;91:317–323.
9. Corbo V, Dalai I, Scardoni M, Barbi S, Beghelli S, Bersani S, Albarello L, Doglioni C, Schott C, Capelli P, Chilosi M, Boninsegna L, Becker KF, Falconi M, Scarpa A. MEN1 in pancreatic endocrine tumors: analysis of gene and protein status in 169 sporadic neoplasms reveals alterations in the vast majority of cases. *Endocr Relat Cancer.* 2010;17:771–783.
10. Swarts DR, Claessen SM, Jonkers YM, van Suylen RJ, Dingemans AM, de Herder WW, de Krijger RR, Smit EF, Thunnissen FB, Seldenrijk CA, Vink A, Perren A, Ramaekers FC, Speel EJ. Deletions of 11q22.3-q25 are associated with atypical lung carcinoids and poor clinical outcome. *Am J Pathol.* 2011;179:1129–1137.
11. Swarts DR, Henfling ME, Van Neste L, van Suylen RJ, Dingemans AM, Dinjens WN, Haesevoets A, Rudelius M, Thunnissen E, Volante M, Van Criekinge W, Van Engeland M, Ramaekers FC, Speel EJ. CD44 and OTP are strong prognostic markers for pulmonary carcinoids. *Clin Cancer Res.* 2013;19:2197–2207.