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***c-MYC* amplification and c-myc protein expression in pancreatic acinar cell carcinomas. New insights into the molecular signature of these rare cancers**

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ABSTRACT

The molecular alterations of pancreatic acinar cell carcinomas (ACCs) and mixed acinar-neuroendocrine carcinomas (MANECs) are not completely understood and the possible role of *c-MYC* amplification in tumor development, progression, and prognosis is not known. We have investigated *c-MYC* gene amplification in a series of 35 ACCs and 4 MANECs to evaluate its frequency and a possible prognostic role. Gene amplification was investigated using interphasic FISH analysis simultaneously hybridizing *c-MYC* and the centromere of chromosome 8 probes. Protein expression was immunohistochemically investigated using a specific monoclonal anti-*c-myc* antibody. 20 cases had clones with different polysomies of chromosome 8 in absence of *c-MYC* amplification, 5 cases had one amplified clone and other clones with chromosome 8 polysomy, while the remaining 14 cases were diploid for chromosome 8 and lacked *c-MYC* amplification. All MANECs showed *c-MYC* amplification and/or polysomy which were observed in 54% pure ACCs. Six cases (15.3%) showed nuclear immunoreactivity for *c-myc* but only 4/39 cases showed simultaneous *c-MYC* amplification/polysomy and nuclear protein expression. *c-myc* immunoreactivity as well as *c-MYC* amplification and/or chromosome 8 polysomy were not statistically associated with prognosis. Our study demonstrates that a subset of ACCs show *c-MYC* alterations including gene amplification and chromosome 8 polysomy. Although they are not associated with a different prognostic signature, the fact that these alterations are present in all MANECs suggests a role in the acinar-neuroendocrine differentiation possibly involved in the pathogenesis of MANECs.

Key words: acinar cell carcinoma; pancreas; MANEC, MiNEN, *c-myc*; amplification, prognosis

INTRODUCTION

Acinar cell carcinomas (ACCs) of the pancreas are a heterogeneous group of cancers showing different morphological features and clinical presentations [1]. In addition, about 10-20% of cases showing a significant neuroendocrine component (> 30% of the tumor burden) are defined mixed acinar-neuroendocrine carcinomas (MANECs). Together with mixed ductal-neuroendocrine carcinomas, they belong to the group of mixed neuroendocrine/non-neuroendocrine neoplasms (MiNENs) [2, 3]. Approximately 50% of ACCs/MANECs are metastatic at the time of diagnosis and about 40% of cases recur as local and/or metastatic disease after surgical resection [4, 5]. The prognosis is poor with 5-year survival rates ranging between 25 and 50% without difference between ACCs and MANECs [5]. Although several attempts have been made to search for morphological, immunohistochemical, and molecular prognostic factors, tumor stage still seems to be the best prognosticator in resectable cases [5].

Our knowledge on molecular alterations of ACCs has been greatly expanded in the last years and several molecular alterations involved in tumor development and progression have been recently identified [1]. Some of them are typical of ACCs, like alterations in the APC/ β -catenin pathway and fusion in *RAF* genes [6, 7], while others involve genes (i.e. *TP53* and *BRCA2*) which play a crucial role in the development and progression of a wide spectrum of different cancers [1]. Interestingly, recently published data have suggested that some molecular alterations, like those involving *TP53*, may identify ACC subtypes showing a more aggressive behavior [8].

It has been recently reported that about 17% of ACCs show *c-MYC* amplification, but the prognostic role of this alteration remains to be clarified [9]. The oncogene *c-MYC* is a transcription factor implicated in about one-third of human malignancies by promoting tumor growth increasing DNA replication and transcription, protein synthesis, cellular metabolism and proliferation [10]. *c-MYC* overexpression is frequently associated with poor clinical outcome [11]

and it has been demonstrated that *c-MYC* plays a pivotal role in the molecular mechanisms underlying the aggressiveness of pancreatic ductal adenocarcinoma [10, 11].

In the present study, we have investigated *c-MYC* gene amplification in a series of 39 pancreatic ACCs/MANECs in order to evaluate its frequency and a possible prognostic role.

MATERIALS AND METHODS

Cases

Thirty-nine cases were selected from our previously-reported series of 62 well characterized pancreatic ACCs [5]. Tumor selection mainly depended on the availability of sufficient material to perform immunohistochemical and FISH analyses together with complete clinical information. The main clinico-pathologic characteristics are summarized in Table 1. All tissues were fixed in buffered formalin (formaldehyde 4% w/v and acetate buffer 0.05M) and routinely processed to paraffin wax.

Immunohistochemistry

For immunohistochemistry, 3µm-thick sections were mounted on poly-L-lysine coated slides, deparaffinized and hydrated through graded alcohols to water. After endogenous peroxidase activity inhibition, performed by dipping sections in 3% hydrogen peroxide for 10 minutes, incubation with primary rabbit monoclonal anti-c-myc antibody (Y69, Abcam, Cambridge, UK) was carried out at 4°C for 18–20 hours, followed by the avidin-biotin complex (ABC) procedure. Immunoreactions were developed using 0.03% 3,3'-diaminobenzidine tetrahydrochloride and then sections were counterstained with Harris' hematoxylin.

Fluorescence in situ hybridization (FISH)

Interphasic FISH analysis was performed on 3-4 μ m-thick sections used for conventional histologic examination as reported in the guidelines of the European Cytogeneticists Association [13] and the cytogenetic interpretation of data agrees with the International System for human Cytogenetic Nomenclature (ISCN) [14]. FISH analysis was performed using direct viewing on a standard fluorescence microscope at $\times 100$ magnification. FISH results were evaluated on representative areas of each tumor identified on hematoxylin and eosin stained slides. To ensure a representative sample and to permit an assessment of the extent of tumor heterogeneity, *c-MYC* amplification and chromosome 8 polysomies were scored in more than 200 interphasic nuclei from at least 5 to 8 separate areas of each tumor by two independent operators (BB and MGT). Only experiments with 90% hybridization efficiency were considered. *c-MYC* amplification was investigated simultaneously hybridizing *c-MYC* (red signal) and the centromere of chromosome 8 (green signal) probes (Zytolight SPEC MYC/CEN8 Dual Color Probe, Zytovision GmbH, Bremerhaven, Germany). Cases were defined as amplified when the ratio (R) between red (*c-MYC*) and green (CEN8) signals was >2.0 . Cases were defined as polysomic for chromosome 8 when at least 20% of neoplastic cells showed three or more copies of CEN8 signals (green signals).

Statistical analysis

Comparisons of continuous data were performed using Student's t tests and discrete variables were compared with χ^2 test or Fisher's exact test. Univariate survival analysis was performed using Kaplan–Meier curves and log-rank test. Data were statistically analyzed using MedCalc® Version 12.5.0.0 and GraphPad Prism Version 5.00 softwares and p value <0.05 was considered significant.

RESULTS

ACCs and MANECs were more frequently observed in males (29 cases) than in females (10 cases) and the average age at diagnosis was 59.7 years (range 33-84 years). Tumors were more frequently located in the pancreatic head (15 cases) followed by the tail (12 cases) and the body region (12 cases). The mean size was 8 cm with a range between 1.6 and 29 cm. 35 out of 39 (89%) cases were pure ACCs, while four cases showing a neuroendocrine cell population >30% were defined as mixed acinar-neuroendocrine carcinomas (MANECs) [3]. The mean follow-up time was 33 months (range 6-135 months). Twenty-seven patients died of disease after a mean follow-up time of 18.6 months, while 11 patients were alive at the last follow-up control (mean follow-up time of 67.5 months). One patient was lost to follow-up.

Six cases (15.3%) showed nuclear immunoreactivity for *c-myc*, in a cell population ranging from 10 to 80% neoplastic cells (Figure 1).

All 39 cases were scored for both *c-MYC* amplification (Figure 2A) and chromosome 8 polysomy (Figure 2B): in detail, 20 ACCs had clones with different polysomies of chromosome 8 in absence of *c-MYC* amplification, 5 cases had one amplified clone and other clones with chromosome 8 polysomy. The remaining 14 cases were diploid for chromosome 8 and lacked *c-MYC* amplification. FISH data of five *c-MYC* amplified cases are reported in Table 2. The clones with *c-MYC* amplification ranged from 33.1 to 77.4% of neoplastic cells. The ratio of *c-MYC* and chromosome 8 centromere ranged from 2.22 to 2.87 indicating presence of low level of *c-MYC* amplification in all cases. Polysomic cases showed different level of polysomies ranging from 3 to 10 chromosomes 8 resulting in gain of MYC copies. Figure 2B shows a polysomic ACC showing nuclei with 6 to 10 chromosome 8. Interestingly, all MANECs showed *c-MYC* amplification and/or polysomy, which were observed in 19 out 35 (54%) pure ACCs. Four out the 39 cases investigated showed simultaneous *c-MYC* amplification/polysomy and a nuclear protein expression. As in most

cases of mixed acinar-neuroendocrine carcinomas [3, 5], the two components were not clearly separated or identifiable on morphological analysis and their identification was performed using immunohistochemistry.

c-myc immunoreactivity as well as *c-MYC* amplification and/or chromosome 8 polysomy were not statistically associated with prognosis ($p=0.04$) (Figure 3).

DISCUSSION

The molecular signature of ACCs is different from that of pancreatic ductal adenocarcinomas and neuroendocrine neoplasms, and more frequently includes alterations in the APC/ β -catenin pathway, while gene alterations frequently involved in ductal adenocarcinomas like mutations in *KRAS*, *DPC4*, *p16*, are absent or very rarely present [1, 15]. Alterations of *TP53* gene (mutation of one allele and loss of the other allele) were recently found to be associated with worse survival [8] suggesting the possibility that specific subtypes of ACC may show specific molecular features with prognostic relevance. In this context, the recent identification that a subset of ACC shows *c-MYC* amplification [9] has suggested us to explore the prognostic role of *c-MYC* alterations in pancreatic ACCs.

c-MYC, whose function is tightly controlled by growth factor-dependent signals in normal adult cells [16, 17], plays a pivotal role in organogenesis and, in particular, in pancreatic acinar cell development and maturation [18]. However, its expression can be deregulated and enhanced via multiple mechanisms in tumor cells and is implicated in the pathogenesis, progression and aggressiveness of several human tumors. In particular, *c-MYC* protein expression and *c-MYC* gene activation by amplification have been found to be associated with tumor aggressiveness and poor prognosis of several cancers [11, 12, 16, 19, 20] including pancreatic ductal adenocarcinoma (PDAC) [10, 11]. Most of genetic and epigenetic alterations playing a role in the pathogenesis and

progression of PDACs involve *c-MYC* activations. *c-MYC* overexpression occurs in about 40% of advance PDACs, although comprehensive genetic analysis demonstrated that *c-MYC* is generally amplified at low levels [21]. Mechanisms involved in *c-MYC* deregulation in PDACs include genetic events or transcriptional, post-transcriptional signaling, or post-translational mechanisms. Genetic aberrations include mutation of the TGF β -inhibitory elements controlling *c-MYC* promoter or *c-MYC* amplification. Transcriptional mechanisms include the activation of transcription factors inducing *c-MYC* transcription or enhancement of *c-MYC* transcriptional elongation by CDK9-mediated phosphorylation of RNA-polymerase. Alterations in post-transcriptional signaling include the attenuation of *c-MYC*-inhibiting miRNAs in absence of *TP53* functions; post-translational mechanisms include CK2-mediated phosphorylation of *c-MYC*, which prevents proteasome degradation resulting in the reduction of *c-MYC* ubiquitination and degradation [10].

In 20 cases of our series we found chromosome 8 polysomy in absence of *c-MYC* amplification and in five cases both *c-MYC* amplification and chromosome 8 polysomy. This result suggests that *c-MYC* activation by gene amplification and/or polysomy is involved in the pathogenesis of at least a subset of ACCs. Interestingly, we found that all MANECs showed *c-MYC* amplification and/or polysomy which, on the contrary, were observed in only 54% of pure ACCs. In this context, it is interesting to recall that *c-MYC* has been found to regulate neuroendocrine trans-differentiation of prostate adenocarcinoma resulting in the formation of the aggressive neuroendocrine-differentiated subtype [22-25]. In addition, *c-MYC* has been demonstrated to play a pivotal role in regulating ductal-neuroendocrine plasticity of pancreatic ductal adenocarcinoma leading to a neuroendocrine differentiation, which contributes to poor outcome and therapeutic resistance [26]. Taking together these findings may suggest that activation of *c-MYC* may lead to an acinar-neuroendocrine differentiation responsible of MANEC development. Starting from this observation, further studies are needed to confirm this hypothesis.

In general, we have found low level of *c-MYC* amplification in our series and this may explain the discordance observed between FISH and immunohistochemistry considering that the latter is a less sensitive method. Immunohistochemical expression of MYC seems to predict well *c-MYC* alterations when more than 50% of nuclei are MYC positive [27]. In our series, among the five cases immunoreactive for MYC only two showed more than 50% of positive nuclei and both cases showed *c-MYC* polysomy.

Regarding a possible prognostic role of *c-MYC* alterations, it is worth noting that pancreatic ACCs and MANECs are a group of aggressive carcinomas showing poor prognosis with 5-year survival rates ranging between 25 and 50%. To date, tumor stage seems to be the only prognosticator in resectable cases [5]. However, among resected cases the search for prognostic factors useful to stratify patients in different prognostic categories is a hot topic in pancreatic pathology. To the best of our knowledge, there are no prognostic factors for surgical resected pancreatic ACCs and MANECs, although in recent years several attempts have been made to search for them. One of our aims was to check whether *c-MYC* amplification could be used as prognostic marker. Although we have observed a trend of worse survival in patients with *c-MYC* amplification than in patients without it we did not find a statistical meaning.

In conclusion, our study demonstrates that a subset of ACCs show *c-MYC* alterations including gene amplification and chromosome 8 polysomy. Although they are not associated with a different prognostic signature, the fact that these alterations are present in all MANECs suggests a role in the acinar-neuroendocrine differentiations possibly involved in the pathogenesis of MANECs.

Author contributions: Design and conception: SLR, BB, MGT; Data gathering and analysis: SLR, BB, MGT, AV, LZ, KN, LA, PB, AS, FS; Manuscript preparation: SLR, MGT, FS.

All authors have read and approved the final version of this manuscript.

Compliance with ethical standard: This study was performed according to the clinical standards of the 1975 and 1983 Declaration of Helsinki and was approved by the Ethical Committee of the Ospedale di Circolo (ASST Sette Laghi), Varese, Italy (n.1277/10).

Conflict of interest: The authors declare that they have not conflict of interest.

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FIGURE LEGENDS

Figure 1. c-myc nuclear immunoreactivity in the majority of neoplastic cells of a pancreatic acinar cell carcinoma.

Figure 2. FISH analysis using MYC probe (red signals) and chromosome 8 centromere (green signals). A subgroup of acinar cell carcinomas and all mixed acinar-neuroendocrine carcinomas showed *c-MYC* amplification (A) and/or chromosome 8 polysomy (B).

Figure 3. C-myc protein expression (A) was not statistically associated with prognosis as well as *c-MYC* amplification (B), chromosome 8 polysomy (C) or their combination (D).

Table 1. Clinico-pathologic and FISH features of the 39 ACCs/MANECs investigated

Case	Sex	Age	Type	Site	Size (cm)	Follow-up (months)	MTS	MYC protein expression	c-MYC amplification	Chr. 8 polysomy*
1	M	49	ACC	B/T	6,2	DOD (24)	no	0	no	no
2	M	51	ACC	H	7,5	DOD (6)	yes	0	no	22%
3	M	57	ACC	B/T	14	AWD (56)	no	0	no	no
4	M	44	ACC	B/T	13	AWD (36)	no	0	no	50%
5	M	70	ACC	B	5	AFD (135)	no	0	R=2.87	25%
6	M	70	ACC	B/T	25	DOD (6)	yes	15	R=2.66	38%
7	M	74	ACC	T	4	AFD (89)	no	0	no	no
8	F	75	ACC	T	3,8	DOD (16)	no	0	no	42.5%
9	M	63	ACC	H	3	DOD (14)	no	0	no	54.5%
10	M	69	MANEC	H	7	DOD (9)	no	0	R=2.22	54.8
11	F	37	MANEC	B	6,2	AFD (111)	no	10	no	66%
12	M	71	ACC	B/T	7	AFD (84)	no	0	no	no
13	M	47	ACC	T	6	AFD (84)	no	0	no	97%
14	F	63	ACC	T	10	DOD (12)	yes	10	no	no
15	M	76	ACC	H	4	DOD (36)	yes	0	no	no
16	M	49	ACC	H	1,6	DOD (79)	no	0	no	no
17	M	55	MANEC	H	5	DOD (6)	yes	0	R=2.83	45%
18	M	67	ACC	T	29	DOD (6)	yes	80	no	22.3%
19	M	67	ACC	T	3	DOD (6)	yes	0	no	no
20	M	71	ACC	H	5	DOD (26)	yes	0	no	100%
21	M	62	ACC	H	4,5	AFD (24)	no	0	no	no
22	M	53	ACC	H	NA	L	yes	0	no	no
23	F	42	ACC	B/T	10	DOD (22)	yes	0	no	93%
24	M	60	ACC	H	5	AFD (76)	no	0	no	100%
25	F	45	ACC	H	11,5	DOD (16)	no	0	no	no
26	M	70	ACC	B	8	DOD (19)	no	0	no	20.6
27	M	47	ACC	T	10	DOD (10)	yes	0	no	15%
28	M	59	ACC	H	3	DOD (22)	yes	0	R=2.77	no
29	M	53	ACC	T	2,5	DOD (8)	yes	0	no	no
30	F	69	ACC	B	5	DOD (30)	no	0	no	13.5%
31	F	72	ACC	B/T	8	DOD (20)	no	0	no	77%
32	M	55	ACC	T	2	DOD (34)	yes	30	no	no
33	M	84	ACC	B/T	7	DOD (9)	no	0	no	85%
34	M	60	ACC	H	4	DOD (13)	no	0	no	no
35	M	46	MANEC	T	16	DOD (12)	yes	0	no	41.5%
36	M	82	ACC	T	11	DOD (26)	yes	60	no	36.4%
37	F	50	ACC	T	4	AWD (32)	yes	0	no	100%
38	F	33	ACC	H	8	DOD (16)	no	0	no	62%
39	F	61	ACC	H	3	AFD (16)	no	0	no	77%

MTS: metastasis; F: female; M: male; ACC: acinar cell carcinoma; MANECs: mixed acinar-neuroendocrine carcinoma; AFD: alive free of disease; AWD: alive with disease; DOD: died of disease; L: lost at follow-up; H: head; B: body; T: tail; R: ratio of *c-MYC* and chromosome 8 centromere; *: percentage of polysomic cells; NA: not available.

Table 2. Correlation between *c-MYC* amplification and chromosome 8 polysomy in acinar cell carcinomas.

case	Amplified cells (%)	Ratio MYC/CEN	MYC Mean value of signals	CEN 8 Mean value of signals
5	38.5	2.87	6.11	2.13
6	60.2	2.66	5.49	2.07
10	33.1	2.22	5.09	2.29
17	77.4	2.83	6.74	2.38
28	44.0	2.77	4.68	1.69

CEN: centromere





