

Clinical significance of multifocal primary tumors in small bowel neuroendocrine tumors

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BACKGROUND

- Multifocal primary tumors are reported to occur in up to 30% of cases of small bowel neuroendocrine tumors.
- Existing retrospective studies suggest that patients with multifocal disease are younger at time of diagnosis, have higher likelihood of experiencing carcinoid syndrome, have worse prognosis and/or disease burden at 36 months, and have shorter disease free survival.
- This study aims to contribute to the understanding of multifocal disease by presenting our experience with this phenomenon at a single institution academic medical center.

METHODS

- Retrospective review of medical records of patients undergoing surgical resection of small bowel neuroendocrine tumor and with complete pathology reports between 2012 and 2017 at a single urban, academic medical center with a dedicated center for neuroendocrine tumor management (The Mount Sinai Hospital, NY) was performed.
- Clinico-pathologic features of patients with multifocal disease were compared to those with single primary tumors.

RESULTS

- Data was abstracted from 62 patients.
- Multifocal disease was seen in 39% of patients.
- As detailed in Table 1, patients with multifocal disease were of the same age as patients with single primary at time of diagnosis and surgical intervention.
- Abdominal pain, diarrhea, and carcinoid syndrome were the most common clinical symptoms in both groups.
- Multifocal disease tended to be well differentiated and have lower Ki67 index. The average size of the primary tumor was similar in both groups.
- Overall survival was similar in both groups.

CONCLUSIONS

- The prevalence of multifocal disease was higher at our institution/series than previously reported.
- No differences in the clinico-pathological features between multifocal and single disease were identified; however, the statistical significance of our findings is limited by small sample size.

RESULTS

	Single	Multifocal
Demographics		
n	38 (61%)	24 (39%)
Female	24 (63%)	13 (54%)
Age at Diagnosis (years +/- st dev)	59.8 +/- 10.5	59.5 +/- 12.8
Age at Surgery (years +/- st dev)	60.6 +/- 10.2	60.3 +/- 13.3
Clinical Symptoms		
Weight Loss	16%	33%
Abdominal Pain	42%	58%
Bowel Obstruction	16%	17%
Diarrhea	61%	50%
Carcinoid Syndrome	45%	46%
Carcinoid Heart Disease	8%	8%
Pathology		
Stage		
I	5%	0
II	0	8%
III	37%	25%
IV	58%	66%
Grade		
G1	66%	80%
G2	32%	20%
G3	3%	0
Ki67 Index		
<2%	65%	86%
2-30%	35%	14%
Size of primary (cm +/- st dev)	1.76 + 0.96	2.39 + 1.63
Outcomes		
Overall survival (as of 2017)	31 (82%)	19 (79%)