

Meeting abstract

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## GIST in a reference cancer center in México

Horacio Noe López-Basave\*<sup>1</sup>, Flavia Morales-Vázquez<sup>1</sup>,  
Esmeralda Patricia Ochoa<sup>1</sup>, Juan Manuel Ruiz-Molina<sup>1</sup> and  
German Calderillo-Ruiz<sup>2</sup>

Address: <sup>1</sup>Department of Gastroenterology, Insituto Nacional de Cancerología, Mexico and <sup>2</sup>Department of Medical Oncology, Insituto Nacional de Cancerología, Mexico

Email: Horacio Noe López-Basave\* - [lobohnoe@aol.com](mailto:lobohnoe@aol.com)

\* Corresponding author

from 24<sup>th</sup> Annual Meeting of the National Cancer Institute of Mexico  
Mexico City, Mexico. 14–17 February 2007

Published: 5 February 2007

*BMC Cancer* 2007, **7**(Suppl 1):A51 doi:10.1186/1471-2407-7-S1-A51

This article is available from: <http://www.biomedcentral.com/1471-2407/7/S1/A51>

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### Background

Gastrointestinal stromal tumours (GIST) are rare malignancies characterised by their association with KIT oncogene mutations. Until now, population-based reports of the incidence or survival of kit-confirmed GIST have been rare, and none have originated in México.

### Materials and methods

We reviewed the files in the Instituto Nacional de Cancerología to identify malignant mesenchymal tumours of the digestive tract between 1995 and 2005, and performed c-kit testing in the tumour samples.

### Results

Seventeen cases were found with 88% of GIST localised in the stomach, 5.8% in small intestine, and 5.8% in esophagus. Fifty-eight percent were classified as high risk of an aggressive behaviour and 42% as low or very low risk. Only one patient received treatment with imatinib mesilate and three had radiotherapy. The relative 3-year survival rate was 29.4% for the entire cohort.

### Conclusion

We report the first review of incidence in a referral cancer center in México. The incidence rate is low and comparable with that of cancer registries from Northern America and Europe. Survival was favourable in our pre-imatinib population although it was low in high-risk cases. Prognostic discrimination of the cases with intermediate, low, or very low risk is inadequate, and these categories should

be considered jointly in the future. Our results will help researchers in establishing baseline values against which they can compare in the future, the impact of imatinib and other Kit tyrosine inhibitors on survival.