## supportive care

1496P

ATTITUDES OF PHYSICIANS TOWARD RISK ASSESSMENT AND USE OF GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) AS PRIMARY PROPHYLAXIS (PP) IN PATIENTS (PTS) RECEIVING CHEMOTHERAPY WITH AN INTERMEDIATE RISK OF FEBRILE NEUTROPENIA (FN)

<u>G. Freyer</u><sup>1</sup>, E. Kalinka-Warzocha<sup>2</sup>, K. Syrigos<sup>3</sup>, M. Marinca<sup>4</sup>, G. Tonini<sup>5</sup>, S.L. Ng<sup>6</sup>, Z.W. Wong<sup>7</sup>, A. Salar<sup>8</sup>, G.G. Steger<sup>9</sup>, M. Abdelsalam<sup>10</sup>, L. Decosta<sup>11</sup>, Z. Szabo<sup>12</sup>

<sup>1</sup>Department of Medical Oncology, CHU de Lyon, Lyon, FRANCE <sup>2</sup>Chemotherapy Department, Wojewodzki Szpital Specjalistyczny im. M. Kopernika, Lodz, POLAND

<sup>3</sup>Oncology Unit, Sotiria General Hospital, Athens School of Medicine, Athens,

 $^4$ Department of Oncology, Grigore T. Popa University of Medicine and Pharmacy lasi, lasi, ROMANIA

<sup>5</sup>Department of Oncology, University Campus Bio-Medico of Rome, Rome, ITALY

<sup>6</sup>Oncology Unit, Bendigo Health Care Group, Bendigo, VIC, AUSTRALIA <sup>7</sup>Oncology Unit, Goulburn Valley Health, The University of Melbourne, Melbourne, VIC. AUSTRALIA

<sup>8</sup>Department of Clinical Hematology, Hospital del Mar, Barcelona, SPAIN <sup>9</sup>Department of Medicine I/oncology, Medical University of Vienna, Vienna,

AUSTRIA

10 Medical Oncology Department, Moncton City Hospital, Moncton, NB, CANADA

<sup>11</sup>Biostatistics, Amgen Ltd, Uxbridge, UK

<sup>12</sup>Medical Affairs, Amgen (Europe) GmbH, Zug, SWITZERLAND

Aim: For chemotherapy regimens with intermediate FN risk (10-20%), physicians should assess patient risk factors to determine whether overall risk is >20% and G-CSF PP is indicated  $\hat{d}^1$ . This study described factors considered important by physicians when assessing FN risk and deciding to use G-CSF PP in pts receiving intermediate-FN-risk chemotherapy

Methods: This prospective observational study (NCT01813721) was conducted in Europe, Australia and Canada. Before pt enrolment, investigators reported their own FN risk threshold at which they usually give G-CSF PP, and selected and ranked factors they considered important when assessing overall FN risk and deciding to give G-CSF PP (investigator baseline assessment). For each enrolled pt, their overall FN risk score and the same factors were assessed before starting chemotherapy (patient assessment), and whether G-CSF PP was planned was reported.

Results: The final analysis included 165 investigators (67% medical oncologists) and 944 pts (median age 61 years, range 20–94) with breast cancer (42%), lung cancer (39%) or NHL (19%). Stage IV disease was reported in 34% of pts; 1% had a history of FN. Table 1 lists factors most often ranked for FN risk assessment and G-CSF PP decision. The median investigator-reported FN risk threshold was 20% and the pt median overall FN risk score was 18%. G-CSF PP was planned in 82% pts with an overall FN risk score≥the investigator's threshold and in 19% pts with an overall risk score<their threshold.

Conclusions: The most frequently considered factors were chemotherapy regimen for assessing FN risk and FN risk assessment outcome for decision to initiate G-CSF PP. Other factors were selected less consistently at the investigator and pt level. A standardised approach to risk factor assessment may improve guideline adherence and G-CSF use. <sup>1</sup>Aapro et al. (2011) Eur J Cancer;47:8-32

Disclosure: G. Freyer: Consultant for Amgen France; E. Kalinka-Warzocha: Received honoraria from Amgen for clinical trials. Given lectures for Amgen; M. Marinca: Has spoken in speaker bureaus for Amgen Romania; L. Decosta: Employed by Amgen and holds stock; Z. Szabo: Employed by Amgen and holds stock. All other authors have declared no conflicts of interest

Factors considered in overall FN risk assessment, % (95% CI)		Factors considered in G-CSF PP decision, % (95% CI)	
Investigator assessment (n = 165)	Patient assessment (n = 944)	Investigator assessment (n = 165)	Patient assessment (n = 944)
Chemotherapy agents in the backbone 88 (82–93)	Chemotherapy agents in the backbone 93 (88–96)	Outcome of FN risk assessment 89 (83–93)	Outcome of FN risk assessment 79 (71–84)
Prior history of FN 83 (76-89)	Tumour type 72 (61–82)	Age 80 (71–87)	Guidelines 67 (53-79)
Baseline laboratory values 76 (68-83)	Guidelines 62 (48-74)	Baseline laboratory values 74 (64–82)	Treatment intent 67 (59-74)
Age 73 (64–81)	Tumour stage 43 (34-53)	Guidelines 71 (61-78)	Age 51 (43-60)
Prior chemotherapy 71 (62–78)	Age 39 (32–48)	Treatment intent 69 (61–77)	ECOG/Karnofsky performance status 36 (28–45)

© European Society for Medical Oncology 2014. Published by Oxford University Press on behalf of the European Society for Medical Oncology. All rights reserved. For permissions, please email: journals.permissions@oup.com.