## 4.6 About the Munich Cancer Registry, its latest annual report and a list of figures and tables

This chapter describes the work of the Munich Cancer Registry (MCR) and aims to help English speaking colleagues, who are engaged in cancer prevention, to interpret results and compare data.

The third annual report of the MCR presents results of cancer registration in the city of Munich and the surrounding areas. As a special focus, the present annual report provides statistical analyses of oesophagus, stomach and prostate cancer. Nearly 7,600 general practitioners, 90 hospitals with about 500 departments and 437 communities will receive this issue. This report highlights the importance and the usefulness of population-based clinical data for physicians.

The MCR is part of the comprehensive Munich Cancer Centre and started registering patients in 1978. For the first years a few departments of the Ludwig-Maximilians-University and the Technical University Munich collaborated. The number of collaborating members grew continuously. These efforts have led to a world age-standardised incidence rate estimated to be 308/100,000 (237 for females) for Munich in 1998.

Bavaria has a total population of 12 million, the Federal Republic of Germany has 82 million people. The MCR collects cancer data from about 2.3 million people, which equals 2.8% of the German population. In 2002, the catchment area will be enlarged to 3.66 million people. The Bavarian Cancer Registration Law came into force at the beginning of 1998 allowing the MCR to legally process all death certificates from the catchment area. Tab. 8a/b illustrate the observed cancer related mortality for the Munich area. DCO-rate decreased to about 12% in 2000. The reduction of the current DCO-rate can be expected as a result of follow-back-procedures in the near future.

Each collaborating hospital receives a survey for all their treated patients, irrespective of their home address. Therefore the MCR also registers patients from outside the registration area. Since the beginning in 1994, the reports of 13 pathological institutions have been the basis for checking completeness of registration. In this way correctness of incidence rates may be roughly judged.

For data collection, 26 cancer-specific forms (fig. 4) have been developed. Often, copies of medical reports are sent to the registry as well. Co-operating hospitals additionally report on local and regional progressions and the occurrence of metastases. In this way, the course of malignant disease is described.

The MCR produces periodical reports for all co-operating hospitals to keep them informed of their clinical registries. The most common cancers in the region are analysed and the larger hospitals have their clinical results compared anonymously.

Fig./Tab. (Abb./Tab.) Page Aims, general conditions and co-operation status Data flow from the 6 clinical cancer registries in Bavaria to the registry of the whole of Bavaria, 2 divided into notification office (depersonalisation) and registration office (data storage&evaluation), and to the national cancer registry in Berlin 2 3 The diverse interests of a cancer registry 3 What information should be transmitted to a cancer registry, by whom and on which parts of the 6 disease process? 4 Example of an MCR tumour-specific data collection form 7 5 The database and different ways of using the data 8 Results 6 Processed data and information in 2000 13 Age-specific and age-standardised incidence rates for men in the city of Munich in 1998 14 7b Age-specific and age-standardised incidence rates for women in the city of Munich in 1998 16 8a Age-specific and age-standardised mortality rates for men in the Munich region for 1998 to 2000 20 8b Age-specific and age-standardised mortality rates for women in the Munich region 22 for 1998 to 2000 9 Percentage distribution of age-specific mortality for men in Germany in 1999 24 24 10 Percentage distribution of age- and tumour-specific mortality for men in Germany in 1999 Percentage distribution of age-specific mortality for women in Germany in 1999 11 25 12 Percentage distribution of age- and tumour-specific mortality for women in Germany in 1999 25 13 Comparison of cancer mortality between the Munich region in 1998 to 2000 and 26 Germany in 1998 14 Extract from the new official death certificate 28 30 15 Cancer deaths in the 155 towns and villages in the catchment area in 1998 to 2000 16 Mapping cancer mortality at a local level in the MCR catchment area 33 17 Breast cancer: Incidence, mortality and proportion of pTis or pT1 patients in the 15 largest 34 towns in the catchment area 18 Characteristics of the most common cancers 38 S1: Number of patients with "good follow-up" and a single malignancy S2-S6: Age at diagnosis S2-S4: 10%, 50% (median), 90% of patients are younger than the indicated age at diagnosis S5-S6: Mean age for men and women at diagnosis S7: Percentage of new male cases S8-S9: Median life expectancy (in years) from diagnosis for cured men and women S10-S11: The official number of tumour related deaths in Germany in 1998 for males (109 thousand) and females (104 thousand) S12-S15: Relative survival rate for 2, 5, and 10 years S16-S17: 5- and 10-year overall survival (all causes of death) S18: 5-year prevalence as a factor of incidence estimated from the overall survival curve S19-S20: Cumulative incidence of secondary malignancies (Kaplan-Meier estimation) S21: Percentage of patients (relative to S1) with at least a second malignancy, who were registered in addition to S1 (underestimated because of underreporting) S22-S23: Number of person years and median follow-up time S24-S27: Percentage distribution of pT findings since 1988 S28: Proportion of lymphnode positive findings S29: Percentage patients with M1 at diagnosis S30-S31: Time (in months) to 1st progression for M0 patients (50%, 90% quantiles) S32-S33: Time (in months) from 1st progression to death (50%, 90% quantiles) S34-S36: Survival time (in months) for M0 patients with at least one progression event (50%, 90% S36-S37: Mean survival time for M0 patients and all patients (incl. M1) with established progression S38: Number of M0 patients with a progression recorded during the disease process S39-S42: Percentage of metastases, local recurrences, regional lymphnode recurrences and unspecified (for solid tumours) progressions during the course of disease

unspecified progression, lung, liver, bone, distant lymph node, CNS and unspecified metastases)

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