The Danish Head and Neck Cancer database

Jens Overgaard1
Aleksandar Jovanovic1
Christian Godballe2,3
Jesper Grau Eriksen3

1Department of Experimental Clinical Oncology, Aarhus University Hospital, Aarhus, 2Department of ORL – Head and Neck Surgery, 3Department of Oncology, Odense University Hospital, Odense, Denmark

Aim of the database: The Danish Head and Neck Cancer database is a nationwide clinical quality database that contains prospective data collected since the early 1960s. The overall aim of this study was to describe the outcome of the national strategy for multidisciplinary treatment of head and neck cancer in Denmark and to create a basis for clinical trials.

Study population: The study population consisted of all Danish patients referred for treatment of squamous cell carcinoma of the larynx, pharynx, oral cavity, or neck nodes from unknown primary or any histopathological type (except lymphoma) of cancer in the nasal sinuses, salivary glands, or thyroid gland (corresponding to the International Classification of Diseases, tenth revision, classifications C.01–C.11, C.30–C.32, C.73, and C.80).

Main variables: The main variables used in the study were symptoms and the duration of the symptoms; etiological factors; pretreatment and diagnostic evaluation, including tumor–node–metastasis classification, imaging, histopathology, and laboratory tests; primary treatment with semidetailed information of radiotherapy, surgery, and medical treatment; follow-up registration of tumor status and side effects; registration of relapse and treatment thereof; and registration of death and cause of death.

Main results: Data from 33,000 patients have been recorded during a period of 45 years. In this period, the outcome of treatment improved substantially, partly due to better treatment as a result of a series of continuous clinical trials and subsequent implementation in national guidelines. The database has furthermore been used to describe the effect of reduced waiting time, changed epidemiology, and influence of comorbidity and socioeconomic parameters.

Conclusion: Half a century of registration of head and neck cancer treatment and outcome has created the basis for understanding and has substantially contributed to improve the treatment of head and neck cancer at both national and international levels.

Keywords: head and neck cancer, squamous cell carcinoma, radiotherapy, national population cohort

About the Danish Head and Neck Cancer group

The Danish Head and Neck Cancer (DAHANCA) group was established in 1976 as a working group by the Danish Society for Head and Neck Oncology with the primary aim to develop national guidelines for the treatment of head and neck cancer in Denmark. DAHANCA group’s activities have included creating and running a national database, development of treatment guidelines, performing numerous national clinical protocol studies (>30), and providing quality assurance programs. The group is multidisciplinary and consists of representatives from all the oncology centers in Denmark treating head and neck cancer, including oncologists, otolaryngologists, pathologists,
computers were introduced in health care). This program later in the radiation physics department was among the first places where a program for the registration of head and neck cancer (the hospital, Aarhus, Denmark, where the medical physicist database at the Department of Oncology, Aarhus University an annual increase of 1,400 patients (Table 1).

The registration of patients with head and neck cancer is the essential backbone of the DAHANCA structure, and the DAHANCA database includes currently >33,000 patients, with an annual increase of 1,400 patients (Table 1).

DAHANCA group has been an active and well-established structure for 40 years and has provided a solid foundation for the treatment of head and neck cancer in Denmark and has become recognized internationally as one of the major head and neck cancer groups. In this aspect, it should be noted that the treatment of head and neck cancer in Denmark has rather a good outcome seen with international eyes.

DAHANCA database
The registration of patients with head and neck cancer is the essential backbone of the DAHANCA structure, and the associated database includes currently >33,000 patients, with an annual increase of 1,400 patients (Table 1).

Historically, the database started as a local home-designed database at the Department of Oncology, Aarhus University Hospital, Aarhus, Denmark, where the medical physicist Mogens Hjelm Hansen in the early 1970s created an elaborate program for the registration of head and neck cancer (the radiation physics department was among the first places where computers were introduced in health care). This program later formed the basis for equivalent databases in Copenhagen and Odense and a subsequent retrospectively registration of, especially, laryngeal cancer back to the early 1960s. The registration of cases later expanded to include the new oncology centers in Aalborg and Herlev, and a full national registration of larynx cancer has taken place since 1971. In the same time period, there have also been regional and more sporadic registrations of pharynx and oral cavity cancers. At a national level, the three initial databases included all patients with larynx cancer between 1971 and 1991.

In connection with a large clinical trial,1 the national database was reconstructed and expanded to include all cancers of the larynx, pharynx, and oral cavity. Each of the five oncology centers had their own databases (named DA 1–5), which were all copies of a central master. Thus, data were gathered locally and were immediately available for daily use and research purposes at the individual institutions. The DA 1–5 data files were regularly merged into the joint database at the secretariat in Aarhus and thus formed the national DAHANCA clinical database. The structure was based on the “MedLog” program. Also, the initial three databases named (AKH, HLM, and HL3) were merged and restructured using the same program. The registration in DA 1–5 covered the period 1992–1999. In 1995, a special DATHYRCA database for registering thyroid cancer2 and more lately databases for unknown primary neck tumor,3 salivary gland tumor,4 and sinonasal cancer were established.5

In 2000, the DAHANCA database was changed into a central web-based database (OCX), which allowed interactive registration directly on a central server with the immediate possibility for quality checks of the data and with instant correction of errors. This increased the quality assurance and gave direct access to information about the current status of the patients. The OCX database is based on the development tool Clarion and uses Hypertext Transfer Protocol Secure certificate with an Hypertext Transfer Protocol Secure connection; all communications are securely encrypted. All users’ logins are logged and registered. The operating system is on a Windows Server 2012 platform, and the server is situated at Aarhus University. The system is flexible and can be modified in connection with specific protocols and projects. All projects are carried out in accordance with the applicable legislation and regulations, including the European Union’s Clinical Trials Directive, and corresponding regulations for the use of good clinical practice and quality assurance, and the database fulfills the Danish Data Protection Agency’s description of standard specification and requirements for security. The OCX database has been active since 2000, and the structure has formed the basis for several other clinical
quality cancer databases (eg, melanoma, sarcoma, ocular cancer, childhood cancer databases). Moreover, in 2011, the DAHANCA database was transformed from being a “private” research database to a clinical quality database in public auspices under the Danish Clinical Registries (RKKP), while still allowing the continued use for research purposes. This simplified the correlation with other public databases, such as the Danish Cancer Registry (DCR), and allowed the database to be used as an indicator of quality.

**Structure of the database**

Approximately 33,000 patients are included in the database, which has ~300 basic variables. Additionally, there is the possibility of specially registered parameters in connection with specific projects. The aim is that established variables remain in the database, and to have, to a large extent, continuous variables that remain unchanged. International standardized classifications are used in this context where such classifications exist (eg, Union for International Cancer Control tumor–node–metastasis classification). With regard to side effects, international principles are also applied, but the principles that we initially developed have now been adopted by other organizations, and thus, the DAHANCA database has contributed to the international standard with regard to registration of side effects from radiotherapy. As not all variables are in use at any time, the daily registration probably involves 100–150 different parameters.

The basic registration includes the following (refer forms at [www.dahanca.dk](http://www.dahanca.dk)):

1. **On study**: symptoms and the duration of the symptoms, etiological factors, pretreatment and diagnostic evaluation, including tumor–node–metastasis classification, imaging, histopathology, and laboratory tests,
2. **Primary treatment**: semi-detailed information of radiotherapy, surgery, and medical treatment.
3. **Follow-up**: registration of vital and tumor status and acute and late side effects.
4. **Recurrence**: registration of relapse and treatment thereof.
5. **Death form**: registration of death and cause of death.

Besides comparison with the DCR, vital status is checked regularly through updates from the civil registration number register. In certain situations, there have been correlations with specific extracts from the National Patient Discharge Registry and the Cause of Death Register.

**How do we register data?**

The principle of the DAHANCA database has throughout the years been a uniform structure of data description and a local registration. For the first many years, no actual clinical record forms existed, but the patient’s journal was considered to be the paper-related data documentation and was used as a basis for data entry. For practical reasons, paper registration forms have been used since 1991 (refer forms at [www.dahanca.dk](http://www.dahanca.dk)). The gathering and recording of data has always been done by a few knowledgeable people, mainly senior specialist doctors. Owing to the limited amount of such people, who have been involved over many years and who know the problems in detail, both communication and collection of data are done in a very uncomplicated and pragmatic fashion. This dependency on a few highly specialized persons is both the strength and weakness of the database, because it not only gives it extremely high quality but also makes it vulnerable to changes. The database has a number of error indicators built in, which safeguards against logical errors and at the same time has the option of reminders if information is missing. Furthermore, lists of irregularities are instantly produced in a continuous attempt to collect missing data or correct any errors.

### Table 1

<table>
<thead>
<tr>
<th>Tumor site</th>
<th>Time period and database acronym</th>
<th>1992–1999 (DA 1–5)</th>
<th>2000–2015 (OCX)</th>
<th>All included</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pre 1971 (AKH, HLM, HL3)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Larynx</td>
<td>857</td>
<td>2,018</td>
<td>10,661</td>
<td>10,661 (9,804)</td>
</tr>
<tr>
<td>Pharynx</td>
<td>254</td>
<td>1,756</td>
<td>10,511</td>
<td>10,511 (8,060)</td>
</tr>
<tr>
<td>Oral cavity</td>
<td>1,304</td>
<td>3,867</td>
<td>5,664</td>
<td>5,664 (5,664)</td>
</tr>
<tr>
<td>Sino-nasal</td>
<td>512</td>
<td>512</td>
<td>902</td>
<td>902 (512)</td>
</tr>
<tr>
<td>Salivary glands</td>
<td>1,697</td>
<td>1,697</td>
<td>5,664</td>
<td>5,664 (5,664)</td>
</tr>
<tr>
<td>Thyroid</td>
<td>1,697</td>
<td>1,697</td>
<td>5,664</td>
<td>5,664 (5,664)</td>
</tr>
<tr>
<td>Unknown primary</td>
<td>1,697</td>
<td>1,697</td>
<td>5,664</td>
<td>5,664 (5,664)</td>
</tr>
<tr>
<td>All</td>
<td>1,225</td>
<td>6,784</td>
<td>19,005</td>
<td>33,768 (29,881)</td>
</tr>
</tbody>
</table>

**Note:** Numbers in bold indicate the time periods with full national coverage of patient cohorts.

**Abbreviation:** DAHANCA, Danish Head and Neck Cancer.
Quality assurance and correlation with other registers

The database is verified through the numerous related research projects. Simultaneously, a specific comparison to the DCR is performed at least annually, where all cases in the DCR have been sought to be verified through the DAHANCA database, and in case of errors (in the DCR), these are explored when possible. Thus, there is currently a full national registration of DAHANCA database ranging back to 1971 (Table 1). This has shown that the DAHANCA database contains 99.7% of the originally known cases in the DCR, and an additional number of patients, currently 2%, have been identified, who were not in the DCR. Furthermore, a number of erroneous registrations in the DCR have been corrected (in the DAHANCA database). The complete interaction between the two databases clearly shows that, in principle, there exists a total nationwide coverage in the DAHANCA database and further ascertains that the DAHANCA database has the potential of increasing the quality of the DCR’s data. The practical aspects of such data feedback to DCR are currently being explored.

This investigative high-quality work has been performed in great detail, and in cases of doubt, relevant additional hospital records and/or other material have been obtained. Thus, all patients in the database have had their hospital records scrutinized at least once, and all updates are verified by a specialist doctor. With the DCR’s well-known almost complete coverage of cancer cases in Denmark as a starting point, it is our impression that with the DAHANCA database we have a near-perfect clinical register, which probably is the world’s most accurate recording of head and neck cancer treatment.

Approval of the database

The initial Aarhus-derived database was approved with register regulations at Vejle County. Later, the national database was registered and approved as a research database by the Danish Data Protection Agency (record nos 1995-1200-282 and 2004-41-4802). Since 2011, it has been a public clinical quality database (record no 1-16-02-249-14). Jens Overgaard is the overall data responsible head, and regionally, the responsibility lies with the consultant responsible for the head and neck cancer treatment at the respective oncological centers.

Economy of the database

As with other research-based databases, the project has initially been based on “con amore” activities performed by interested professionals at weekends and in late hours. Gradually, the need for organized help has increased and the database has throughout the years been partly financed in connection with research projects, where among others, PhD students were assigned to the activities mainly sponsored through the Danish Cancer Society and the Medical Research Council (we have constantly tried to avoid any commercial influence and sponsoring). Since 2006, we have been partly supported thorough the DMCG/RKKP structure. However, the activities have throughout benefited from being part of the infrastructure at the Department of Experimental Clinical Oncology at Aarhus University Hospital and from the numerous dedicated persons who have used their spare time to strengthen the database.

Use of the database

The value of any database is linked with its use, and the quality data are accessible through access to the RKKP (http://www.rkkp.dk/forskningsadgang/). Throughout the years, the DAHANCA database has been extensively explored and has served as an active research database that has formed the basis for numerous scientific projects and publications, including 15 PhD and doctoral theses (for full list, refer the DAHANCA homepage [www.dahanca.dk]). It has been the supporting database for >30 prospective clinical trials and has acted as a baseline for large retrospective studies describing the natural history and treatment outcome in head and neck cancer. Examples of studies are given in Figure 1, and more can be found on the DAHANCA web site. This means that the database is constantly revised and updated and, consequently, is extremely well verified.

Since many of the DAHANCA protocols have focused on biological-based improvement of head and neck cancer treatment, these translational studies have been based on a large amount of clinical material. This DAHANCA-related biobank contains biological material (tumor, normal tissue cell lines, blood samples, DNA, RNA, etc) from >4,000 patients with detailed clinical data and follow-up. Also, paraffin-embedded tumor biopsies are available from almost all patients. Currently, we are systematically collecting material from new patients in the National Danish Cancer Biobank.

The database has not only been strengthened by linkage to other registries but has also shown its value by being a supportive basis for studies of comorbidity, value of follow-up, or socioeconomical relationship in patients with head and neck cancer. In the foreseeable future, it will be part of international comparisons of treatment strategies, and for such purpose, a clinical database is far superior than using cancer registry comparisons (such as the Nordic NORDCAN study), since such databases are incomplete and inaccurate.
Regarding clinical information. Thus, it is important to use the right tools/databases for the right purposes, and when conclusions about therapeutic strategies are the aim, the data must be obtained from a well-verified clinical quality database. We believe that the DAHANCA database is a prime example of such an activity.

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Disclosure
The authors report no conflicts of interest in this work.

References


