In the course of the development of the *NeuroDiseaseMonitor*, the German Center for Neurodegenerative Diseases in Bonn (DZNE) and the Cologne, organized the workshop "**Monitoring Dementia – Towards a Dementia Monitoring System and a Dementia Registry in Germany**" November 17-18, 2011. The aim of the workshop was to discuss the use of registry and health insurance data for the purposes of dementia monitoring.

The workshop brought together experts who have either established a national dementia registry, or are working with registry or health data in the area of dementia and cognitive functioning. The following aspects were discussed:

- Research findings about the prevalence and incidence of dementia in Europe
- Existing population-based data sources that permit epidemiological studies
- The need for new data sources
- Population-based data from health registries and health insurers
- The contribution of these data to a better understanding of the risk of dementia
- The validity of data from health insurers for the purposes of monitoring dementia
- The creation of a national dementia registry in Germany
- Lessons from the experience of existing dementia registries, as well as national cancer registries

The presentations can be accessed here:

- After 50 years of community studies what do we know about dementia? (Michael E. Dewey)
- SveDem Swedish Dementia Registry - a national clinical quality database on dementia disorders (Maria Eriksdotter)
- Registries in Germany: Establishment and use of registries based on routine health data (Jürgen Stausberg)
- Interest and Limits of the French Alzheimer's registry (Jean François Dartigues)
- Patients with dementia: What do we learn from claims data? (Ingrid Schubert, Veronika Lappe)
- Dementia and long-term care (Heinz Rothgang)
- The Diagnosis of Dementia in Public Health Insurance Data (Gabriele Doblhammer, Anne Schulz)
- Utilization of health routine data for outcomes research (Christian Günster)
- Longstanding studies in or run from Cambridge Institute of Public Health (Carol Brayne)
Workshop 2013

The DZNE aims at strengthening the interdisciplinary scientific exchange "Neurodegenerative Diseases: New Insights from Registries, Cohorts and Health Care Data" by using registers, population-based cohort studies, as well as administrative data.

Detailed information about the program can be found here: