Estimating the Burden of Cancer

Projections of the incidence of cancer serve several purposes. These range from making investment decisions on cancer treatment facilities and planning manpower requirements, to formulation and evaluation of policies for control of cancer.

Different statistical methods can be used for such projections. In recent years, these have been translated to user-friendly software application modules. The methods include the simple linear regression method (LR),1 the join point regression method (JP),² and the age, period and cohort model (APC).³ The first two methods can be used with or without taking age as a factor. When age is not taken into account, the crude incidence rate (CR) is used. For all these methods, the primary requirement is reliable data on incidence for various cancers for at least a sample of the population. A well run population-based cancer registry provides valid incidence and mortality rates for the defined populations that they cater to. Since 1982, the National Cancer Registry Programme (NCRP) of the Indian Council of Medical Research has been collecting such information through its population-based cancer registries which now number 27. Though these registries cover only 7% of the population of India, they do provide a fair idea of the magnitude and patterns of cancer, which allows reasonable extrapolation of the burden of cancer for the rest of the country. The publication of the report on 'Time trends in cancer incidence rates: 1982–2005' further strengthened the accuracy of such estimates on the burden of cancer.

The APC modelling for projection takes care of non-linear trends in the data including the possible contribution of period and cohort to the trend. However, one needs to show whether the latter two actually have an effect so as to alter the predictions in a major way, especially in the Indian context. The Surveillance, Epidemiology, End Results Program (SEER) in the USA uses the JP regression method to study changes in time trends and to provide estimates of various cancers.

We predicted cancer cases by all the above methods—LR with CR, LR with effect of age, JP with CR, JP with effect of age and APC modelling—using the incidence data of Bengaluru (for 1982–2001) and Delhi (for 1988–2002) to project the burden of cancer for 2006–07. We then compared the projections by different methods with the actual data reported for 2006–07. The difference in the numbers obtained by different methods was also examined keeping the APC model as the base (Table I).

The estimated number of cases for both Bengaluru and Delhi, using the populationbased cancer registry data with all 5 methods including APC modelling, were lower

Table I. Estimated number of cases (≥25 years) for specific sites by 5 different methods for Bangalore* and Delhi† (2006–07)

Method		City		Site of cancer							
			Men			Women					
			All sites	Prostate	Colon	All sites	Breast	Cervix	Ovary		
LR	CR	Bangalore Delhi	4430 12 265	251 806	142 315	5374 11 066	1377 3211	703 1382	307 920		
	Age	Bangalore Delhi	3997 11 959	256 819	149 267	5172 10 992	1389 3084	684 1336	290 898		
JP	CR	Bangalore Delhi	3713 11 594	254 831	122 321	5129 11 119	1422 3220	786 1534	249 822		
	Age	Bangalore Delhi	4073 11 418	235 667	127 268	4932 10 887	1450 3256	822 1519	382 866		
Age period cohort (APC)		Bangalore Delhi	4362 10 540	243 747	112 292	5213 10 464	1432 2997	849 1661	338 867		
Based or NCRP R		Bangalore Delhi	5349 12 318	372 959	216 295	6760 11 905	1867 3370	1066 1826	380 896		

^{*} based on population-based cancer registry data of 1982–2001 † based on population-based cancer registry data of 1988–2002 LR linear regression JP join point CR crude rate NCRP National Cancer Registry Programme Values that are significantly (p<0.05) lower than that derived through the APC model are shown in bold while those significantly higher are underlined

than the actual reported data. This was almost uniform among men and women except for all-sites among women using the JP and CR method and ovarian cancer using the LR and CR as well as LR and age method for Delhi where the numbers were slightly higher than those for the APC modelling estimates. The prediction for both Bengaluru and Delhi using the LR and CR method were comparable to those observed with the APC method, again with some exceptions such as for cancer of the cervix where it was significantly lower and for all-sites in men and women in Delhi where it was significantly higher. Thus, the different methods gave different estimates though most methods erred in providing lower estimates.

It is essential to adopt a systematic scientific and statistical approach to estimating prediction of number of cancer cases for various anatomical sites of cancer, especially in a developing country such as India. The paucity of data from the rural areas other than the northeast region makes this task even more difficult. The article by Swaminathan *et al.*⁶ in this issue provides one standard method to estimate the burden of cancer in Tamil Nadu based on data from Chennai city and Dindigul district (a predominantly rural district). Using a combination of the data, the paper provides estimates using APC modelling for the state of Tamil Nadu. The authors acknowledge that their estimates could err on either side—more or less than the actual numbers. Nonetheless, the availability of these estimates would be useful for policy-makers and planners.

Since the commencement of the NCRP in 1982, annual, bi-annual and consolidated reports have been regularly published. A brief report on time trends in incidence rates was presented in the consolidated report for 1990–96.7 The first systematic report on trends in incidence rates over 2 decades⁴ shows a steady and consistent increase in the age-adjusted incidence rates of certain cancers across all major urban registries. Among men, cancers of the prostate, colon, rectum and liver have shown statistically significant increase in incidence. Cancer of the prostate is the leading site of cancer among men in most western countries as is cancer of the colon. Among women, cancers of the breast, corpus uteri and lung have shown a rise. While the first two of these cancers could be accounted for because of cohorts with later age at marriage, decreasing multi-parity and so on, the increase in lung cancer could be attributed to an increase in the use of tobacco by women. Lung cancer in women may also be increasing because of environmental exposure to smoke (passive smoking). Three other sites of cancer that have shown an increase in incidence rates in women are ovary, thyroid and gallbladder. The increase in gallbladder cancer is seen in registries that have recorded a comparatively lower incidence than Delhi, which showed an increasing trend only during the earlier years, with a decline in more recent years. There have been rising incidence rates for cancers of the brain as well as in tumours of the lymphoid and haemopoetic system, especially non-Hodgkin lymphoma in both men and women. The decline in the incidence of cancer cervix is seen across all registries including the rural registry at Barshi. This decline is observed in the absence of any organized screening or early detection programmes in the registry areas. The factors contributing to an increase in breast cancer could possibly be responsible for the decline in the incidence of cancer of the cervix. Another possible reason for the decline could be an increase in the number of child-births at institutions (as opposed to home deliveries) leading to improved maternal and maternity care including genital hygiene. This could be a result of the family welfare drive initiated by the government about 4 decades ago and which is continuing. Better genital hygiene, barrier contraceptive use and superior nourishment could all have contributed to the reducing incidence of cancer of the cervix.

The NCRP data that have accrued over the years are essentially that of selected urban centres and only one rural registry that covers part of a district is represented. Therefore, it would be difficult to provide valid estimates of the burden of cancer for the entire country with over 70% of the population of India residing in rural areas. Nonetheless, limited exercises have been done and the paper by Swaminathan *et al.* is another step towards obtaining more projections of cancer in India. Table II provides a summary of estimated new cancers from the first systematic report on trends in incidence rates over 2 decades for India.⁴

TABLE II. Estimated new cancers at all anatomical sites (ICD-10: C00-C96)

Year Men Women Total 2008 447 399 498 773 946 172 2009 454 842 507 990 962 832 2010 462 408 517 378 979 787 2015 497 081 563 808 1 060 889 2020 534 354 614 404 1 148 758				
2009 454 842 507 990 962 832 2010 462 408 517 378 979 787 2015 497 081 563 808 1 060 889	Year	Men	Women	Total
2010 462 408 517 378 979 787 2015 497 081 563 808 1 060 889	2008	447 399	498 773	946 172
2015 497 081 563 808 1 060 889	2009	454 842	507 990	962 832
	2010	462 408	517 378	979 787
2020 534 354 614 404 1 148 758	2015	497 081	563 808	1 060 889
	2020	534 354	614 404	1 148 758

REFERENCES

- 1 Boyle P, Parkin DM. Cancer registration: Principles and methods. Statistical methods for registries. IARC Sci Publ 1991;95:126–58.
 2 Kim HJ, Fay MP, Feuer EJ, Midthune DN. Permutation tests for join point regression with applications to cancer rates.
- Stat Med 2000;19:335–51. Erratum in Stat Med 2001;20:655.

 3 Parkin DM, Muir CS, Whelan SL, Gao YT, FerlayJ, Powell JJ (eds). Cancer incidence in five continents, Vol. VI. Lyon,
- France:IARC Sci Publ No. 120; 1992.
- 4 National Cancer Registry Programme (NCRP, ICMR). Time trends in cancer incidence rates (1982-2005). Bengaluru: NCRP-ICMR; 2009.
- 5 National Cancer Registry Programme (NCRP, ICMR). Three-year report of population based cancer registries (2006-8). Bengaluru:NCRP-ICMR; 2010.
- 6 Swaminathan R, Shanta V, Ferlay J, Balasubramanian S, Bray F, Sankaranaryanan R. Trends in cancer incidence in Chennai city (1982–2006) and statewide predictions of future burden in Tamil Nadu (2007–16). Natl Med J India
- 7 National Cancer Registry Programme (NCRP, ICMR). Consolidated report of the population based cancer registries (1990-1996). Bengaluru:NCRP-ICMR; 2001.

T. RAMNATH A. NANDAKUMAR Coordinating Unit National Cancer Registry Programme (ICMR) Nirmal Bhawan, ICMR Complex Kannamangala Post Bengaluru Karnataka

ank@blr.vsnl.net.in, ncrpblr@canceratlasindia.org