NEWSLETTER SAMPANDAN



QUATERLY NEWSLETTER

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Preservation of neurocognition in the era of high precision radiotherapy - a short review

Uday Krishna and Rakesh Jalali

Introduction

The anatomic and pathologic features of invasion of most the glial tumours is a limitation for complete surgical excision. Radiotherapy hence remains a vital and most effective adjunct to surgery in the CNS malignancies. In a landmark publication in 1978 by Walker et al of the Brain tumour study group¹ confirmed a survival benefit of radiotherapy in malignant gliomas. A systematic review by Laperriere et al has clearly shown that post-operative external beam radiotherapy is recommended as standard therapy for patients with malignant glioma. In low grade gliomas, the EORTC, RTOG and other studies have shown clear role of radiotherapy in delay of the time to recurrence, symptom control and survival benefit.

Poor penetration of chemotherapeutic agents of the blood brain barrier makes radiotherapy the ideal modality of treatment in brain metastasis from systemic malignancy. Addition of prophylactic cranial radiotherapy as an adjunct to systemic chemotherapy has demonstrated survival benefit in patients with small cell and non small cell lung cancer and in children with leukemia. Late effects in the form of endocrine dysfunction, neurocognitive deficits, and second malignant neoplasms have been attributed to RT. The biology of neuro cognition, various theories, effects of RT on cognition and the literature demonstrating role of RT in neurocognitive decline have been discussed.

Biology of neuro cognition and pathogenesis of radiation induced damage

Declarative memory (explicit memory/ remembering) is defined as the capacity for conscious recollections of facts and events. The structures of medial temporal lobe are the hippocampus (including the dentate gyrus) and adjacent anatomically related cortical areas, the entorhinal, perirhinal and parahippocampal cortices. The perceptual processing in neocortex gets transmitted to the medial temporal lobe structures and persists as long-term memory. The Projections from neocortex arrive initially in the parahippocampal cortex and perirhinal cortex and later to the entorhinal cortex and the hippocampal formation, thereby relating these structures get access to ongoing cortical activity. Diencephalon and mammillary bodies also receive inputs from the hippocampus. The medial temporal lobe and medial thalamus, thus constitute the neural system essential for the formation of long-term memory².

In the hippocampus, memory function is associated with principal cells which includes pyramidal and granule cells of the dentate gyrus. New granule cells formed from the stem cells in the sub granular zone, migrate to the granular cell layer, develop granule cell morphology, neuronal markers and connect to their target area. Cranial irradiation can lead to significant injury to normal brain structures. These effects are although imminent at higher doses of radiation, cognitive dysfunction without a structural damage is seen at low doses, in both adult and paediatric population. Cognitive decline manifests as deficit in memory, learning and spatial information processing².

Monje et al at the UCSF have demonstrated this phenomenon of "stem cell hypotheses" in mice. Apoptosis of the hippocampus was assessed immunohistochemically using bromodeoxyuridine (BrdUrd), 6- 48 h after whole brain radiotherapy³. Apoptosis was high 12 h after irradiation and was dose dependent. The proliferating sub granular zone was reduced to 96% at 48 h after irradiation and immature neurons were decreased from 40 to 60% in a dosedependent fashion. The long term consequences were quantified with neurogenesis 2 months after irradiation with 0, 2, 5, or 10 Gy. The production of new neurons was significantly reduced by irradiation, being dose dependent with no effects on the production of new astrocytes or oligodendrocytes.

Effects of Whole brain radiotherapy on neurocognition

Poor penetration of blood brain barrier by chemotherapy agents and better control of microscopic disease by radiation has defined the role of WBRT in local control of patients with brain metastasis and survival in patients with lung cancer and childhood leukaemia.

From AROI, ICRO office & AROI Newsletter Editorial Board

With the hustle and bustle of general Loksabha elections now settled, we have a new ruling party at the helm of national affairs with a lot of expectations from the electorate. We look forward to a new direction and strong commitment from Government in healthcare sector particularly cancer care.

If you look at our events calendar, academic activities have already started and will peak around last quarter of 2014 at National & International level. We must prepare ourselves and participate whole heartedly by showcasing our activities at academic and professional front.

ICRO, ASTRO and ESTRO collaborative courses are a great platform for young radiation oncologists to update themselves and they must make use of such opportunities actively.

Best wishes always,

Dr. Rajesh Vashistha Secretary General AROI

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Preservation of neurocognition in the era of high precision radiotherapy - a short review

Chang et al through a phase III randomised study at the MD Andersen cancer centre, showed neurocognitive decline at 4 months in patients who received WBRT after stereotactic radiosurgery (SRS) compared to those patients who received SRS alone.⁴ Radiation induced neuronal injury to hippocampus and medial temporal lobe was considered central to neuro cognitive decline.

RTOG 0933 is a prospective phase II study which compared the decline in neurocognitive function by hippocampus sparing WBRT using tomotherapy based IMRT with the historic controls who had received WBRT without sparing hippocampus, as a part of PCI studies for SCLC. Previously, the group had demonstrated that the peri hippocampal region (hippocampus plus 5mm margin) risk of recurrence of brain metastasis was 8.6%. RTOG 0933 showed better preservation of delayed recall as assessed by Hopkins verbal learning test (HVLT-DR) without compromise on local control in patients who underwent hippocampus sparing IMRT compared to the historic controls⁵. At 4 months post RT, decline in the HVLT- DR was 7% compared to 30 % in the historic controls (p=0.003) and only 2% of patients undergoing 6 month post RT evaluation showed further decline.

Temporal lobe and hippocampus dosimetry and correlation with decline in neurocognition for partial brain radiotherapy

A prospective controlled observation study by Gondi et al in adult patients with benign or low grade brain tumours treated with fractionated stereotactic radiotherapy and observed a dose response relationship between radiation dose to hippocampal dentate gyrus and long term memory impairment. A dose greater than 7.3 Gy to 40% of the bilateral hippocampi was associated with long term impairment of delayed verbal recall as measured by Wechsler memory scale⁶.

Correlation of dose delivered to the left temporal lobe and decline in neurocognition was initially demonstrated by Jalali et al, initial results of a large randomised controlled trial comparing efficacy of stereotactic conformal radiotherapy (SCRT) versus conventional RT in children and young adults with low grade brain tumours. Neurocognitive decline defined as a >10% drop in baseline intelligence quotient (IQ) correlated with doses more than 43.2 Gy to 13% of the left temporal lobe⁷.

Radiation dose levels on the hippocampus and various other brain structures was analysed by Jalali et al to preserve neurocognition in young patients with low grade brain tumours treated with high precision stereotactic conformal radiotherapy (SCRT). Change in the intelligence quotient (IQ) after SCRT was correlated with dose to bilateral hippocampi and other normal brain structures. Comparison of dosimetric data of patients with >10% drop in baseline IQ with the patients who maintained a normal IQ revealed that in patients receiving >27Gy (50% of the prescribed dose) to >50% volume of the left temporal lobe, showed an IQ decline⁸.

Modifications in neuraxis irradiation in medulloblastoma and neurocognition

Merchant et al have shown that children with high risk (HR) MB, requiring higher doses of craniospinal irradiation and posterior fossa boost show greater decline in neuro cognition compared to those with average risk (AR) MB (who receive craniospinal irradiation to a dose of 23.4 Gy and posterior fossa boost to a dose of 36 Gy). However this was not consistent in children less than 7 years of age in whom efforts to reduce the posterior fossa boost dose did not translate to cognition preservation⁹.

Hyper fractionated RT provides an advantage of dose escalation while maintaining the therapeutic ratio in terms of increased biological equivalent dose to the cranio-spinal axis, while reducing the biological equivalent dose to the normal nervous system and minimising late effects. Eventually this translates to a higher local control and better relapse free survival and overall survival. This was tested for AR MB by various groups in the world. HIT-SIOP-PNET4 was large multi institutional randomised trial addressing this issue comparing hyper fractionated RT and conventional RT, and showed that at a median follow up of nearly 6 years, there was no difference in relapse free survival or overall survival. Late effects in terms of ototoxicity were not different in the two arms.

A similar large longitudinal study was conducted by Gupta and Jalali et al. RT was delivered with two daily fractions (1Gy/fraction), 6-8 hours apart and 5 days/week. CS axis received 36Gy/36 fractions, followed by conformal tumour bed boost 32Gy/32 fractions for a total tumour bed boost of 68Gy/68 fractions. Cognitive assessment was prospectively performed pre treatment and specified post treatment follow up visits¹⁰. Early results showed preserved cognition for all tested domains in children evaluable at 2 years after completion of RT with no significant decline over time.

Conclusions

Hippocampus is the seat for processing spatial information along with other structures of the temporal lobe. Clinical studies with SCRT in children and young adults with low grade brain tumours showed greater cognitive decline with higher doses to these structures. With improved local control and survival of patients with systemic cancer metastasising to brain and low risk of recurrence in the peri hippocampal region, techniques to spare hippocampus during WBRT (such a tomotherapy) to reduce cognitive decline are very promising. Exploring the radiobiological superiority of hyper fractionation in treating the entire craniospinal axis with IMRT (such as tomotherapy) in children with average risk medulloblastoma is proving to be useful. Optimal utility of technological refinement in radiation planning and delivery is necessary to reduce late toxicity like neuro cognition in cranial irradiation.

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Gupta T, Jalali R et al. Early clinical outcomes demonstrate preserved cognitive function in children with average-risk medulloblastoma when treated with hyper fractionated radiation therapy, Int J Radiat Oncol Biol Phys, 2012 1; 83(5): 1534-40.

| S.No. | Name of Fellowship | Nos | For | Age Group (Years) | Fellow- ship Grant (in Rs) | Basis | LM AROI since (Yrs) | Min. Papers | Regularly Attending AROI confer- ences | Already availed fellowship in last 5 years | |
|-------|--|-----|--------------------------|-------------------------|-------------------------------------|---|------------------------------|----------------|--|--|--|
| | Overseas | | | | | | | | | | |
| 1.1 | AROI- Kirloskar Therapeutics | 2 | Radiation Oncologists | >50 | 1.5 lakhs | MD/ DNB | 10 | 5 | yes | No | |
| 1.2 | AROI- DR. Reddy's Lab AROI- Merck India Ltd. AROI- Novartis India Ltd. | 2/3 | Radiation Oncologists | 41-50 | 1.5 Lakhs | MD/DNB | 10 | 5 | yes | No | |
| 1.3 | AROI- Dr. Reddy's Lab AROI- Fulford AROI- Nucleotron | 2/3 | Radiation Oncologists | 35-40 | 1 lakh | MD/DNB | 5 | 5 | yes | No | |
| 2. | Within India | 4 | | | | | | | | | |
| 2.1 | AROI- Dr. Reddy's | 2 | Radiation Oncologists | 30-35 | 50,000 | MD/DNB | 3 | 1 | yes | No | |
| 2.2 | AROI | 1 | Medical Physicist | <40 | 20,000 | DRP/MSc (MP) | 2 | 1 | Yes | No | |
| 2.3 | AROI | 1 | Radiation Oncologists | < 35 | 20,000 | MD/ DNB | 3 | - | - | - | |
| 2.4 | AROI | 1 | RT Tech- nologist | <45 | 10,000 | AERB certified | yes | | | No | |
| 2.5 | Neil Joseph Fellowship | 5 | Radiation Oncologists | 5 | 10,000 | Student MD/DNB 3 rd Year PG | yes | In conf | | | |
| 3 | Best Paper | | | | | | | | | | |
| 3.1 | Best Proferred Paper for Senior Members | 1 | Radiation Oncologists | 40-50 | | Post MD/ DNB >10 Yr | 10-15 | In conf | yes | | |
| 3.2 | Dr. M S Gujral Best Proferred Paper Senior Members | 1 | Radiation Oncologists | 35-40 | | Post MD/ DNB 5-10 yr | 5-10 | In conf | yes | | |
| 3.3 | Dr. G C Pant Young Doctor Award | 1 | Radiation Oncologists | <40 | 20000 + plaque | Post MD/ DNB 3 yrs | 3 | In conf | yes | | |
| 3.4 | Gold Medal Parvati Devi | 1 | Radiation Oncologists | | 10000 + medal | | yes | In conf | | | |
| 3.5 | Gold Medal Dr. J. M. Pinto | 1 | Radiation Oncologists | | 7500 + Medal | | yes | In conf | | | |
| 3.6 | Best paper | 1 | Medical Physicist | <30 | 5000 | DRP/MSc (MP) | yes | In conf | | | |

AROI FELLOWSHIPS / AWARDS

Conditions

1. Applicants have to send a copy of date of birth certificate.

2.Applicants to send a copy of the publications mentioned under each Fellowship

3. Self certified proclaimation that they are working full time in radiotherapy.

4. Fellowship amount will be given to candidates from money received from sponsors after tax deduction and 15% contribution to AROI fund.

- 5. All the applications for fellowship/ best paper awards be sent along with the letter from head of department/ institute to the office of Secretary General AROI by 5 PM, August 31st 2014. 6. No Objection certificate from their head of Department if selected to go for
- fellowship

7. PG Students shall send their certificates through Head of the Department.

8. For the best paper award, applications should be sent along the full paper.

9. Abstract along with the letter from the head of dept. for publication in JCRT should be sent along with the paper.

10. Radiation Oncology Applicants above age 35 must be member of ICRO. 11. Applicants to send softcopy also thru email.

Mailing address:

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16th ICRO PG Teaching Course at Shimla



Proud student s with distinguished faculty

16th ICRO PG teaching course was held on 26-27th April, 2014 The faculty was invited from Apollo hospital Hyderabad, HCG in Shimla and the theme was "Radiotherapy Planning. From Banglore, TMH Mumbai, GCRI Ahmedabad, PGI Chandigarh, 2D to 3D".

There was an overwhelming response and total 93 students from all over the country participated. The program was inau- The program was highly appreciated by not only students but gurated by Prof Jayshree Sharma, Director Medical Education, Himachal Pradesh.

The course was started with lecture on basic physics of radiotherapy planning and continued with radiotherapy planning for head and neck cancers, pelvic tumors, Ca breast, Ca esophagus, Ca lung and cranio-spinal axis irradiation.

PGI Rohtak Fortis Hospital Gurgaon, Medanta Medicity Gurgaon, and IGMC Shimla.

also by consultants. Finally quiz based on MCQs from the various lectures delivered in the course, was conducted and two students stood first. One was from TMH Mumbai and other was from IGMC, Shimla.

Input by Dr. Manoj Gupta

Notice

36th Annual conference of AROI (AROICON14) is going to be held in Imphal (Manipur) from 6th Nov to 9th Nov 2014. Main points of discussion will be as follows:

GBM of ICRO

On 6th Nov, 2014 after the completion of ICRO work shop at same hall.

GBM of AROI - I

On 8th Nov. 2014 at 6 PM or after completion of best paper in the main hall at Imphal

Agenda will be given later on.

Agenda for GBM

Amendment of constitution a. Introduction of the post of "President Elect"

- Change of tenure of President/Secretary b. Post/Editor-in-chief.
- c. Increase in registration fee for new members.

Detailed agenda will be posted on website and mailed to all members as well.

GBM of AROI - II

30 minute after completion of GBM-I

Active participation of members is solicited.

Report from Apollo Cancer Conclave, 2014

Apollo Hospitals Group organized a major international cancer conference-**Apollo Cancer Conclave, 2014** from 6th – 9th **Feb. 2014**, in **Chennai.**

The conference is to commemorate the 20th anniversary of the establishment of the Apollo Oncology Services in India. As cancer care has become one of the fastest growing health care needs in our country, Apollo Group chairman is keen to bring together the best minds in oncology both from India and overseas under one roof to deliberate on the rapid advances in this field and its practical applications.

The focus of this conclave is on integration of the cutting edge science, technology and emerging advances, into the day to day management of the patient care in the Indian Subcontinent, and the South East Asian region.

The Pre Conference Live workshops and Lung cancer Symposium were heldo on 6th Feb 2014 at the Hyatt Regency, Chennai. It included Uro oncology surgeries & Upper GI Oncology prodedures live from Apollo Hospitals Chennai, Head & Neck Live Workshop and the High precision Radiation therapy live workshop from Apollo Speciality hospital. As a unique, the robotic Live Surgery workshop by Dr.Vipul Patel was live transmitted from Orlando, USA. On 7th Feb '14 NeuroSurgery live workshops are webcasted in Apollo Speciality hospital. All the workshops are webcasted in Apollo Cancer Conclave website and we live transmitted to various association and the few institutions.

The academic lecture sessions was started on 7th Feb 2014 at ITC Grand Chola, Chennai. We had 3 parallel tracks for the sessions; We had dedicated Meet the Professor sessions which was structured timetabled form in which 15 registered focused delegates discussed with the International expertise and exchanged their knowledge.

The conclave was inaugurated by the former President of India Dr.APJ Abdul Kalam on 7th Feb'14, Healthy deliberation were done to improve the cancer care in the entire Asian and African region the dignitaries from various countries in detail at Afro Asian Symposium which was held on the same day.



Special Plenary session was held on 8th Feb'14 with the guest of honor Satguru Jaggi Vasudev , Dr.Shantha , Chairman Adyar Cancer Institute & Dr.B.Ajai Kumar, Chairman, HCG cancer hospitals. Women Cancer Symposium and the Conquerors night held on the same day.

Apart from the dedicated scientific sessions we had special sessions on Robotic surgery, High Precision Radiation Therapy, Molecular Biology, Special Plenary sessions, Cancer Screening, women Cancer Symposium and Cancer Conquerors night and Afro Asian Symposium.

The Apollo Cancer Conclave was very successful and had a participation of over 2000 delegates from 28 countries, including 300 faculties with 36 scientific sessions dedicated to each specific area in cancer care.

There was a overwhelming response and the appreciation from all the delegates who really were very satisfied with the high quality academic discussions.

An International Journal has agreed to publish the important outcomes from our Apollo Cancer Conclave. The same can be sent to Govt. of India for the benefit of the medical community of our country.

Obituary

Dr Sanjay Supe (1962 - 2014)

Born in family of illustrious father Medical Physicist Dr. S J Supe at Mumbai, he destined to be a Medical Physicist.

He was jovial, full of life, research oriented person who was friend, philosopher& guide to so many physicists and oncologists.

For almost three decades he made KMIO Bengaluru as his home and made a distinct name for himself in radiation oncology fraternity.

His sudden loss due to illness has created a great void and we pray to almighty for peace to his departed soul.

Some of comments on facebook on his demise

I am shocked, Sanjay was energetic colleague. Fearlessly frank, brutally honest. My heartfelt condolencesto bereaved family.

- K S pathasarthy

Its very sad, we have lost a fine human being. - Suresh Pamidighantam

Call for participation in 36th AROI conference at Imphal from Nov. 06-09, 2014

The "36th Annual Conference of Asso-

ciation of Radiation Oncologists of India" is being organized by North Eastern Chapter of AROI and being hosted by Department of Radiotherapy, Regional Institute of

Medical Sciences, Imphal, Manipur (India) during Nov. 06 - 09, 2014. The theme of the conference is "Collaborate and together we conquer". The first day of the conference will be devoted to CME on brachytherapy and the remaining days will deliberate on all types of cancer.

Conference Venue:

Collaborate and together we conquer—enjoy this themed conference in the lap of Manipur, the Jeweled Land, full of surprises and rich culture

Conference Secretariate:

Department of Radiotherapy,

Regional Institute of Medical Sciences, Imphal. Manipur (India) – 795004 Tel.: 0385-2410901 / Fax: 0385 -2411703 Email: aroicon2014imphal@gmail.com

The conference

will be held in

"City Convention

heart of the city.





Our Newsletter Partners for this issue



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Forthcoming Events 2014

National

July 2014

12-13 17th ICRO Teaching Course Tata Memorial Centre, KOLKATA

Email: raj.shrimali@tmckolkata.com

August 2014

15-16 MhCl AROI Nagpur

Email: mhci.aroi@gmail.com

September 2014

13 4th Teaching Course "Basic Radiobiology" RCC, IGMC, Shimla

Email: mkgupta62@yahoo.co

20-21 20th NZAROICON-2014 PGIMS, Rohtak

> Email: nzaroicon2014@gmail.com secretary@nzaroicon2014.com

26-28 FHNO-2014 PGIME&R, Chandigarh @ J W Mariott

Email: vamahospitality@hotmail.com www.fhno2014.com

October 2014

10-11 12th Annual Radiotherapy Practicum Teaching Course in Radiotherapy Tata Memorial Centre, Mumbai

Email; radonco@tmc.gov.in

AROI members going places

Dr Rakesh Jalali was invited as a **Grand Round Speaker at the MD Anderson Cancer Center**, Houston on 25th November 2013.

The title of his talk was "Pediatric Neuro Oncology: challenges and successes in developing countries".

Dr. Rakesh Jalali Co-chaired the Scientific Committee at the recently held World Federation of Neuro Oncol-

11-12 UPAROI-2014

R.R. Cancer Institute & Research Centre, Bareilly

Email : uparoi2014@srmsims.ac.in Website : www.srmsims.ac.in

19-23 AROI-ESTRO COURSE ON ADVANCED TECHNOLOGIES CANCER INSTITUTE, ADYAR, CHENNAI

Email: vsrinivasan09@gmail.com October 2014

November 2014

20-22 AMPICON-2014

Rural Medical College, Pravara

Email : ampicon2014@gmail.com Website : www.pravara.com/ampicon2014.html

December 2014

26-28 ISMPOCON-2014 Hotel Hyatt Regency, KOLKATA

> Email : ismpocon2014@gmail.com Website : www.ismpo.org/ismpocon

International Events

November 2014

28-30 9th SFO, 2014 The Leela - Ambience, Gurgaon

> Mail:
> saarconcology14@gmail.com, akvaid@yahoo.com
>
>
> Web:
> www.sfo2014.com

> > Compilation by :Dr. Pardeep Garg, Faidkot

ogy (WFNO) conference in San Francisco between 21st to 24th November,

Dr. A K Anand, Director, Radiation Oncology moderated a session on Treatment Planning during "Awareness Programme on Radiation Safety & Quality Assurance: Driving the benefits of Innovations in Radio Diagnostic and Therapy" on June 5, 2014

4. Dr. Subhash Gupta, Astt. Prof. gave a lecture on Evolution of Modern Radiation Technologies during above mentioned session

Fortis Memorial Research Institute holds CME cum Workshop on Electron Therapy-Revisited

FMRI, Gurgaon had a CME cum Workshop on electron application in Radiation Oncology on June 14, 2014. Event covered topics related to physics of electrons, current technologies for its generation, dosimetry, commissioning, beam planning, special therapy procedures and clinical challenges in its application. sessions and give hands on training. A lively debate on whether electron therapy has lost its relevance in modern day therapy was held at the end of day.

A large number of Radiation Oncologists and Medical Physicists participated in this event.

Input by T Ganesh & Kanan Jassal



Eminent faculty from national and international centres participated to address the

Response to last issue question

Role of evidence based Radiation Treatment Management in current practices ?

Evidence based practice is a norm globally and its definition is, "The conscientious ,explicit and judicious use of current best evidence in making decisions about the individual patient care".

It means our professional judgements and behaviours should be guided by two independent principles:

When ever possible practice should be grounded on prior findings that demonstrate empirically...that they are likely to produce predictable, beneficial and effective results.

Every client system over time should be evaluated.

In radiation oncology, current practice is based on Individual's clinical expertise supported by NCCN & RTOG guidelines which helps in multi centre trials and evaluations for systematic research

Compilation of various current Radiation Oncology practices by Dr. Arun Verma, Consultant, Max Pat-

parganj, Delhi indicates effect of evidence on cancer management as follows:

A systemic review and meta-analysis recently published in green journal by Marta et al had established role of IMRT over 3D CRT. It showed significant overall benefit of IMRT in xerostomia score grade 2-4 with insignificant trend of improved local control and survival.

In another study, Radiotherapy in breast cancer patient added a survival benefit of 3.8% in 15 years in post BCS, 5.4% in 5 years in post mastectomy, node positive patients. Though, newer techniques do not add survival benefit but have role in delivering homogenous doses and reducing late toxicity specially cardiac morbidity, maintaining size of breast / cosmetics

Similarly novel technique i.e. SBRT has challenged surgery in T1-2 lung cancer with 95% local control and 66.6% overall survival at 2 year while randomised controlled trial and meta-analysis established chemoradiation as standard of treatment in stage III NSCLC and limited stage SCLC.

INT 0116 trial made adjuvant chemoradiation standard of treatment in loco-regionally advanced gastric cancer with median OS of 35 months vs 27 months in surgery alone.

Systemic review and meta-analysis published in JCO 2012, established role of adjuvant RT+CT in R1 operated hepato-biliary tumors, and uncertain benefit for R0 patient with positive node but in contrast there is level 3 evidence of improved survival benefit in postoperative T2-3, N0-1M0 gall bladder cancer.

Although randomised trial showed equal survival benefit with preop

When ever possible practice should be grounded on prior findings that demonstrate empirically...that they are likely to produce predictable, beneficial and effective results

or postop RT+CT in rectal cancer but there is added advantages associated with preop RT+CT like increased respectability with down staging, and increased pathological response which may get reflected in increased survival benefit with increased sphincter preservation rate in long term and future trial.

> Prostate cancer is one of site where newer technique hypo-fractioned IMRT, IGRT +/brachytherapy challenged surgery in all three risk group i.e. low, intermediate and high risk. Another site is cervical cancer where role of radiotherapy is strongly established and limiting role of surgery in medically fit, stage Ia only. Ib and IIa can be equally treated by surgery or EBRT + Brachytherapy but most of time required adjuvant RT in presence of risk factors. Beyond this RT+CT is gold standard treatment except in IVB.

> Level of evidence, its source, grade of recommendations decide whether guidelines are recommendations, suggestive or no guidelines are possible from the study.

An article recently published in journal 'Report of Practical Oncology and Radiotherapy that explored evidence based radiation oncology and emphasized development of new radiotherapy centre and upgradation of existing centre to improve quality of radiotherapy that get reflected in term of better outcome with good quality of life.

However implementing any new evidence based practice is always a challenge as its relevant advantage, costs, compatibility, complexity, trialability, reinvention, observability of benefits and risk attributes need to be addressed

Input by Dr. Arun Verma & Deepak Arora

Views expressed are personal by respondent to question of the issue.-Editor

Question of this issue

Are we ready for incident free radiation treatment delivery ?

Please send your reply for publication in next issue of AROI newsletter to deepak.arora3@maxhealthcare.com