Expert opinions on Robotic CyberKnife Technology

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ABSTRACT

Robotic Cyber knife (CK) is an advanced robotic Stereotactic Radiosurgery technology that is used in Radiation oncology to treat brain and body tumours in cancer patients. The aim of this study was to find out and compare how Cyber knife Image guided technology is being used in two active clinical Cyber knife centres in USA and Australia to provide treatment to patients with brain and other cancers. The expert opinions of a medical physicist and a Medical Radiation therapist with clinical experience of using Cyber knife were captured using an E-questionnaire. This study assessed clinical, technical, organizational and Educational strategies and resources employed to provide Cyber Knife treatment in two clinically active CK centres. This study was done in 2020.

KEYWORDS: Brain tumours, CyberKnife,Image guided Radiotherapy, Real-time tracking, Stereotactic body Radiotherapy, Stereotactic Radio surgery.

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I. INTRODUCTION

Robotic Photon based CyberKnife Image guided Radiotherapy is currently being used in certain institutes globally to provide Stereotactic radiation treatment for intra and extra cranial tumours. The CyberKnife System is developed by Accuray Incorporated, Sunnyvale, California, USA[1]. The CyberKnife system delivers both radiosurgery (SRS) and frameless stereotactic body Radiotherapy (SBRT). The definition of SBRT is provided by Stereotactic Radiotherapy working group upon the request from German Society of Radiation Oncology and this definition is agreed by other working groups in different countries. According to thisdefinition SBRT is a form of external Beam Radiotherapy that delivers highly conformal high radiation doses in few fraction with image guidance. SBRT also involves active or passive intrafraction motion management and follow up [2].

The CyberKnife System consists of six main components namely i) a 6 MV linear accelerator mounted on ii) a robotic arm and iii) a tumour tracking system (In-room stereoscopic KV x-ray system with in-floor detectors, iv) Respiratory motion management system (Synchrony), v) treatment couch with 5 degree of freedom and vi) an algorithm that connects the tumour motion with chest wall motion in order to predict tumour motion at all times during the treatment[1]. Synchrony system monitors patients' breathing in real time and consists of Infrared Light emitting diodes placed on patient's thorax along with wall mounted infrared detector or camera. It ensures that linear accelerator is synchronized with target that moves due to respiration.

CyberKnife has five tracking options namely 6D Skull, X Sight spine, X Sight lung with Synchrony, Fiducial with Synchrony, and Fiducial [3]. XSight tracking system that is good for spine tumours but is not good for abdominal tumours which are positioned distal to spine [4]. CyberKnife system provides AI driven real time tumour tracking of implanted fiducial markers and respiratory motion management to ensure treatment accuracy by constantly identifying and correcting for tumour and radiation beam mismatches throughout the entire treatment. The use of CyberKnife is increasing globally and it is first SBRT and SRS technology that provides real time tumour tracking. Above all CK treatment is associated with sharp dose fall. These features of CK technology ensures accuracy in treatment delivery and therefore is likely to result in use of reduced treatment margins resulting in better sparing of Organs at risk. This in turn ensures dose escalation resulting in potentially better tumour control and reduced treatment induced side effects. The purpose of the study was to gather opinions of CK experts to find out how Robotic photon based Cyber knife Image guided technology is being used to provide treatment to patients with brain and other cancers. The study wants to identify variations in dose

prescription and margin and tumour tracking methods. The present study discusses how Cyber Knife technology is used in two Institutions based in Australia and USA, what clinical, technical and organization resources are used to impart CK treatment, what challenges were faced during its implementation and what improvements are sought in the CK technology by the experts. The study also recorded what education and training pathways are used to impart CK knowledge. The present study gives a synopsis of similarities and differences in employing CK technology for management of various cancers.

II. MATERIALS AND METHODS

- A. Study OverviewAn expert opinion E-survey was designed to gather opinions and views of Radiation Oncology professionals who have expertise in CyberKnife Treatment Planning, delivery and dosimetry.LinkedIn platform (Social Media) was used to contact experts of CyberKnife technology. Two experts based in CyberKnife centres in Australia and USA agreed to fill in the survey and the E-survey was sent to them via LinkedIn. The study was conducted in 2020.
- **B.** Selection of case studies and E-SurveyThe questionnaire was designed in MS word and consisted of 33 questions, most of which were close ended questions. Survey questions were structured in five sections namely i) Demographic, ii) CK institutional background Information, iii) CK information, iv) Treatment planning and delivery, v) Knowledge and Experience. Appendix A shows sample E-survey.
- **C. Ethical Considerations** This study was deemed IRB exempt as it was a quality enhancement and evaluation study. Responses were anonymous so no ethical approval was required. No patients were approached. No medical or personal data of participants collected. By answering the questionnaires, the professionals agreed to give their informed consent.

D. Statistical Analysis

Data was recorded and analysed in Microsoft Excel. Descriptive analysis was used to examine the results of the study.

III. RESULTS

A. Respondent Characteristics: 100% of the respondents were male. 50% of respondents belonged to 30-40 years of age range and the other 50% belong to 50-70 years of age range. both respondents were married (100%). One respondent was Medical Radiation Therapist from Australia and other was Medical Physicist from USA. Medical Physicist from US has 10 years of Clinical Experience of CK (50%) whereas Medical Radiation Therapist from Australia had 5 years of clinical experience. Results are show in Figures 1-3

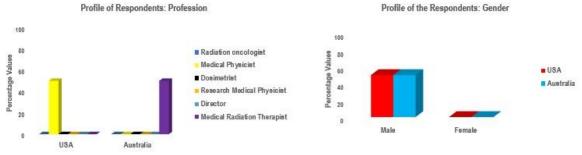


Fig.1 Socio Demographic Profile of Respondents: Gender

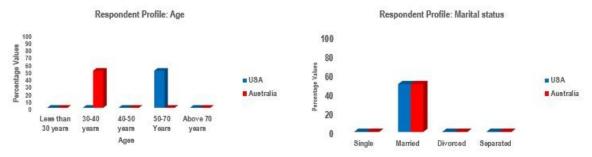


Fig. 2 Socio Demographic Profile of Respondents: Age and Marital status

How long you have been using CK and SBRT to give treatment?

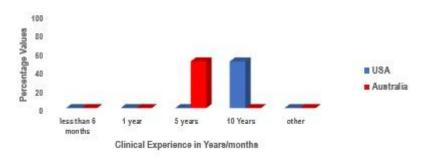


Fig. 3 Clinical Experience of CK

B. CK institutional Background and Resources

1. Do you have CK in your hospital or Institution?

The respondent from USA (50%) said there was no CK in his hospital whereas Respondent from Australia had CK (50%).

2. Location of the Institute and Type of Practice

One respondent 950%) was from Nevada, USA and the other from Perth Australia (50%). Results are shown in Fig 4-5.

Geographic location of the Institution

100 80 80 80 USA Australia

Perth

Fig. 4: Location of CyberKnife Centres

City / State

Reno, Nevada

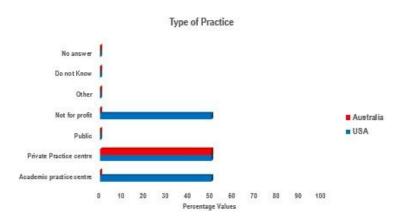


Fig. 5 Type of practice

3. Professionals involved in CK delivery

Respondent from USA said that two medical physicists (13.3%), one radiographer (6.7%), four Radiation therapy Technologist (26.7%), three radiation oncologist (20%) and 5 neurosurgeons (33.3%).

whereas respondent from the Australia only mentioned that medical physicists, radiographers, radiation technologists, dosimetrist andradiation oncologists all are involved in CK treatment planning and delivery but did not specify their number. Results from US case study are shown in Fig. 6.

How many professionals are involved in CK Treatment Planning & Delivery?

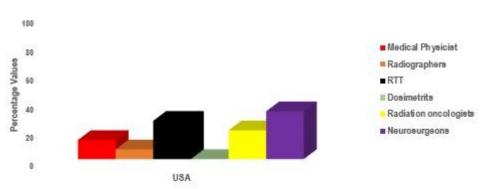


Fig. 6 CyberKnife Team

4. Technical, Clinical & Organizational Challenges

The respondent from Australia said they experienced financial difficulties while respondent from US said they experienced no challenges in implementing CK. Results are shown in Fig.7

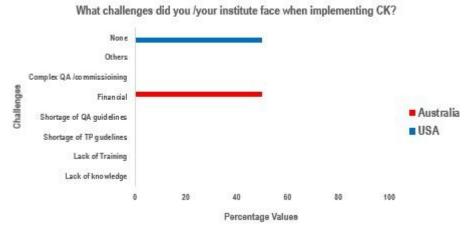


Fig. 7 Challenges

C. Information about CK Technology

1. Intent

The respondent from Australia said CK is used for both Curative and palliative purposes whereas the Medical physicist from US said CK is used for curative purposes. Fig.8 shows the results.

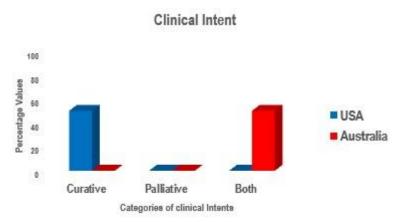


Fig. 8 Clinical Treatment Intent

2. Type of tumours

Both respondents from Australia and US said they use CK to treat both Intracranial and body tumours as well as for primary and metastatic tumours. Results are shown in Fig. 9-10

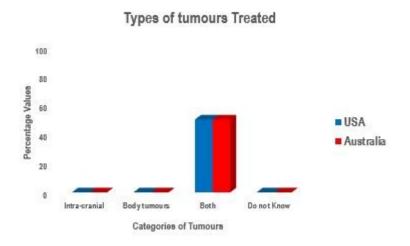


Fig. 9 Tumour Types

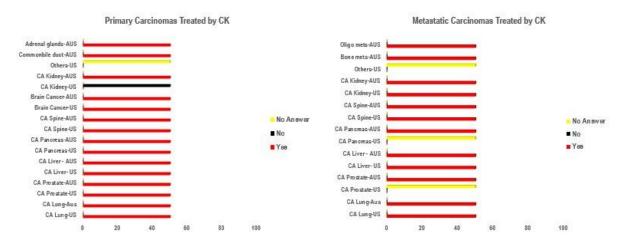


Fig. 10 Primary and Metastatic tumours

3. Stage

Respondent from US said CK is only used for early stage cancers whereas respondents from Australia said CK is used for both Early and advanced stage disease in their hospital. Results are shown in Fig.11

Do you use CK for Early or advanced stage Cancers?

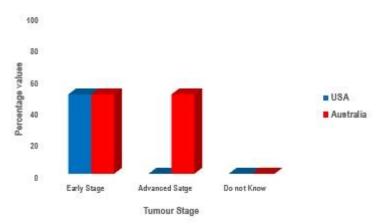


Fig.11 Tumour Stage

4. Reasons for CK adoption in the department/organization

Results are shown in Table I.

| 7 | Γable I |
|----|----------|
| CK | Adoption |

| Cir nuoption | | | | | | |
|----------------------------|----------------------------|--|--|--|--|--|
| Reasons for CK Adoption | | | | | | |
| USA Australia | | | | | | |
| | Dose Escalation | | | | | |
| Precise treatment delivery | Precise treatment delivery | | | | | |
| Better local control rates | Better local control rates | | | | | |
| | Treatment time reduction | | | | | |
| Retreatment | Retreatment | | | | | |
| | Clinical Research | | | | | |
| Gain Competitive edge | Gain Competitive edge | | | | | |

D. Treatment Planning and Delivery

1. CK Image Guidance System

In response to the question what CK image guidance system consists of both respondents mentioned tracking. Results are shown in Table II.

Table II CK components

| Components of CK IGRT | | | | | | |
|-------------------------------------|-------------------------|--|--|--|--|--|
| USA Australia | | | | | | |
| Tracking Using Orthogonal KV system | Skull Tracking | | | | | |
| | Spine Tracking | | | | | |
| | Synchrony with fiducial | | | | | |
| | 1 view lung | | | | | |
| | 2 view Lung | | | | | |
| | Fiducial Tracking | | | | | |

2. Other IGRT Systems

In response to the question what type of Image guidance you use to localize target & verify target before treatment delivery responders stated various IG systems in addition to CK image guidance. Results are shown in Table III

Table III

| Image guidance | | | | | | |
|--|----------------------------|--|--|--|--|--|
| IG used for Target Localization& Verification | | | | | | |
| USA Australia | | | | | | |
| | In room volumetric imaging | | | | | |
| | Planar imaging | | | | | |
| CK image Guidance System | CK Image Guidance System | | | | | |
| Fiducial Marker (Except for brain, spine & most lungs) | Fiducial Markers | | | | | |
| During treatment: Imaging for continuous tracking | | | | | | |

3. Immobilization

Various immobilization devices are used in conjunction with CK treatment. Results are shown in Table IV.

Table IV
Types of Immobilization applied during CK

| Immobilization Techniques/Devices | | | | | | |
|-----------------------------------|------------------|------------|--|--|--|--|
| Cancers | USA | Australia | | | | |
| CA Lung | | | | | | |
| CA Prostate | | | | | | |
| CA Liver (HCC) | | | | | | |
| CA Pancreas | | | | | | |
| Spinal Cancer | Mask for C-spine | | | | | |
| Brain Cancer | Mask | Head Frame | | | | |
| Kidney Cancer | | | | | | |
| Others | | | | | | |

4. Treatment dose and fractionation

Different dose regimes are used in US and Australia. Results are shown in Table V.

Table V
Dose Regimes

| Mo | st Common Dose Regimes | |
|--------------------------------|------------------------|---------------|
| Cancers | USA | Australia |
| Primary Localized PC | 36.25Gy in 5# | 35-36Gy in 5# |
| Metastatic PC | Boost | 35-36Gy in 5# |
| Primary Lung Tumour (ES) | 60Gy in 3 or 5# | 54Gy in 3# |
| Primary Lung Tumour (AS) | | 54Gy in 3# |
| Metastatic Lung cancer | 60Gy in 3 or 5# | 54Gy in 3# |
| Recurrent lung cancer | 50-60Gy in 5# | 54Gy in 3# |
| Primary Unresectable small HCC | No answer | 45Gy in 3# |
| Primary large unresectable HCC | No answer | 45Gy in 3# |
| Liver Metastases | No answer | 54Gy in 3# |
| Recurrent Unresectable HCC | No answer | 54Gy in 3# |
| Primary Spinal lesions | No answer | 27Gy in 3# |
| Metastatic spinal lesions | No answer | 27Gy in 3# |
| Primary Pancreatic lesion | No answer | 40Gy in 5# |
| Metastatic Pancreatic lesion | No answer | 40Gy in 5# |

5. Margins

In response to the question how much margin you apply to GTV to get CTV, respondent from US stated zero CTV margin for listed cancers whereas respondent from Australia stated 2mm margin for both advanced and early stage primary lung carcinomas. Results for both CTV and PTV are shown in Fig. 12-13 and Table VI.

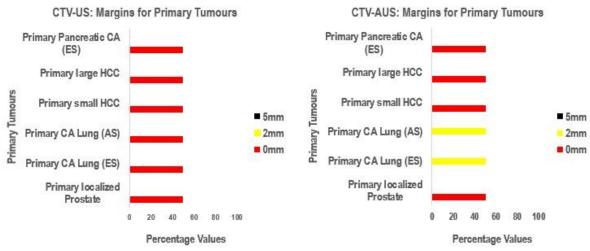


Fig. 12 CTV=GTV+margin

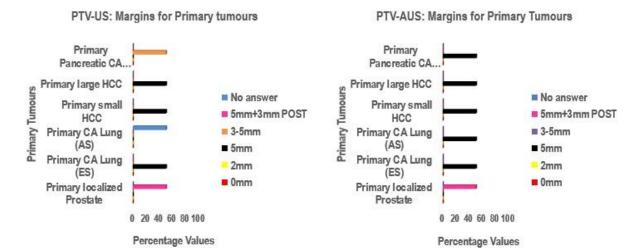


Fig. 13 PTV = CTV + margin

Table VI Margins for Metastatic Disease

| Cancers | USA CTV=GTV + Margin | USA PTV=CTV + Margin | Australia CTV=GTV+ Margin | Australia PTV=CTV+ Margin |
|---------------|-------------------------|-------------------------|------------------------------|------------------------------|
| Spinal | 0 | 1mm | 0 | 0 |
| Metastases | | | | |
| Liver | 0 | 5mm | 0 | 5mm |
| Metastases | | | | |
| Localized | 0 | 3-5mm | 0 | 5mm |
| Pancreatic | | | | |
| cancer (ES) | | | | |
| Metastatic PC | 0 | Boost | 0 | 5mm+3mm post |

6. Beam Energy

Both respondents said they use 6MV for various tumours. Results are shown in Fig. 14

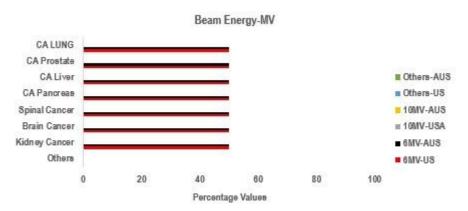


Fig. 14 CK System Beam Energy

7. TPS and Radiation treatment planning

The medical Physicist in US said CyberKnife Multiplan Treatment planning system (50%) is used whereas respondent from Australia said Accuray Precision radiotherapyTPS (50%) is used for CK radiation treatment planning. In Australia Radiation Technologist whereas in US Medical Physicist perform CK radiation treatment planning. Results are shown in Fig. 15-16.

Treatment Planning Systems used for CK

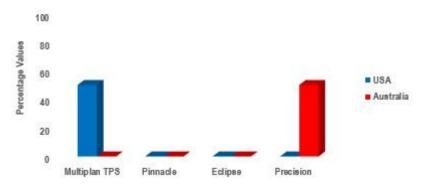


Fig. 15 CK TPS

Who Performs Treatment Planning in your Organization?

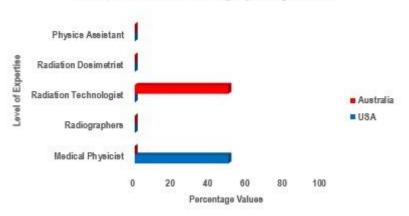


Fig.16 Professions responsible for Treatment planning

8. Motion management

In response to the question which technique is employed to manage breathing induced motion, both respondents (100%) said they use fiducial markers and Synchrony Respiratory tracking system (real time tracking). Results are shown in Table. VII-VIII

Table VII
Motion Management Techniques-USA

| Cancers | USA | USA | USA | USA |
|-------------|-----------------|-------------------------|-------------|-------|
| | Fiducial Marker | Synchrony TM | Breath hold | Other |
| Ca lung | Y | Y | N | - |
| CA prostate | Y | N | N | - |
| CA Pancreas | Y | Y | N | - |
| CA liver | Y | Y | N | - |
| CA Kidney | Y | Y | N | - |

Note: Y= Yes, N=No, CA=carcinoma

Table VIII Motion Management Techniques-Australia

| wiotion wanagement rechniques-Austrana | | | | | | | | |
|--|-----------------|-------------------------|-------------|--------------------|--|--|--|--|
| Cancers | AUS | AUS | AUS | AUS | | | | |
| | Fiducial Marker | Synchrony TM | Breath hold | Real time tracking | | | | |
| Ca lung | Y | Y | | - | | | | |
| CA PC | Y | N | | Y | | | | |
| CA Pancreas | Y | Y | | - | | | | |
| CA liver | Y | Y | | - | | | | |
| CA Kidney | Y | Y | | - | | | | |

Note: Y= Yes, N=No, CA=carcinoma

- 9. **Common toxicities:**respondent from Australia said fatigue and tiredness was most common acute toxicity experienced by patients suffering from prostate, liver, pancreas and kidney cancers. Respondent from USA did not answer the question.
- 10. **CK vs. IMRT vs. VMAT vs. Proton Therapy:**in the view of expert from US, CK treatment provides better tumour control, Disease free survival (DFS), Overall survival (OS) and reduced CK treatment induced morbidity compared to IMRT,VMAT and proton Therapy. The CK Expert from Australia said it is a complex question and cannot be answered.

E. Knowledge and Experience

Results are shown in Fig.17-19

Acquisition of CK Knowledge & experience by staff

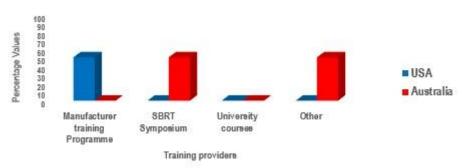


Fig. 17 Pathways for achieving CK Education & Training

Are Radiation oncology/Medical Physics Programmes in your country provide SBRT and CK experience?

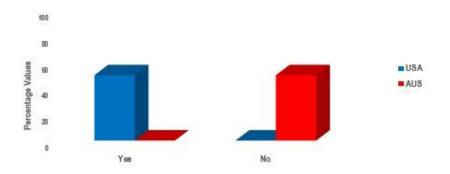


Fig. 18 CK Experience

Is CK treatment cost effective comapred to VMAT, IMRT,

Tomotherapy?

100

80

90

USA

Australia

Fig. 19 CK Cost Effectiveness

IV. DISCUSSION

Discussion

The present study provides a description of current practices of Robotic CK as well as clinical, technical and organizational resources used in imparting CK treatment for treating both intracranial and extracranial tumours in Australia and USA.

The present study is unique as it compares the CK practices, resources and strategies via Expert opinions in two institutions (case studies) based in widely different geographical regions. The present study collated the data on Profession, Gender, age, Marital status and Clinical experience of CK experts under the category of demographic Information.

Both CK experts had significant clinical experience of using CK (Australian Expert: 5 years, USA Expert: 10 years). The respondent from Australia was Medical Radiation Therapist based in a private hospital in Perth while respondent from the US was a Medical Physicist currently working in a private not for profit academic centre in Nevada. He previously worked in another centre that had CK facility.

Consensus differed between the two experts with regards to Challenges faced during CK implementation in the department, number and types of professionals required for CK treatment planning and delivery, Intent of CK treatment, stage of the disease, RT dose and fractionation. In the present study, the respondent from Australia said they experienced financial difficulties while respondent from US said they experienced no challenges in implementing CK. A paper by Dieterich and Pawlicki [5] highlights the complexity of QA program for CyberKnife in clinical practice and recommend that frequency of QA checks should be based on clinical studies rather than on historical benchmarks established for massively different technologies. The study also recommends formation of phantoms appropriate for distinctive QA needs of CyberKnife system.

In terms of similarities of CK practices, CK technology is used to treat both intracranial and extracranial tumours. CK is also used to treat both primary and metastatic disease in both case studies. However there was some differences. Expert from USA said CK is used to treat primary lung, prostate, liver, pancreas, spinal and brain tumour but not used to treat primary kidney tumours in his centre. He also said metastatic tumours of lung, liver, spine, brain and kidney are treated by CK. Expert from Australia said that CK is used in his centre to treat both primary and metastatic tumours of Lung, prostate, liver, pancreas, spine, brain, kidney, CBD (Common bile duct), adrenal tumours. In addition to it in Australia CK is used to treat bone metastasis and oligo metastasis. Literature review also shows that CK is treated for various primary and metastatic tumours [6–8].

In the present study Different doses were reported for lung cancers by both experts (60 Gy in 3-5# in USA vs. 54Gy in 3# Australia). Most common dose was spinal cord tumours was 27Gy in 3 fractions, for liver tumours 45 Gy - 54 Gy in 3 fractions, for prostate cancer was 35-36Gy in 5 fractions and for pancreatic tumours 40Gy in 5 fractions. In the present study fatigue and tiredness were most common acute toxicity experienced by patients suffering from prostate, liver, pancreas and kidney cancers.

In the present study no margin was added to GTV to obtain CTV for localized prostate disease and a margin of 3mm posteriorly and 5mm anteriorly and laterally was added to obtain PTV. This is in line with the literature. Both respondents from US and Australia reported using 3-5 mm margin around GTV to obtain PTV for localized pancreatic cancer. This is similar with the margins used in Song et al. [9] study who expanded GTV by 3mm to get PTV. In the present case studies a margin of 2mm was added to achieve PTV for spinal metastases in USA case study and a margin of zero in Australian case study. A margin of 5mm was reported in both case studies for liver metastasis. These margins are similar to margins reported by Kato et al. [10] in liver cancer patients.

Dose Regimes, Margins and Radiation induced Toxicity: NSCLC studies

There is 0% risk of developing radiation myelitis when treated with hypo-fractionated regimes of 8Gy in 1 fraction to 4Gy in 5 fractions [11-12]. A review of three randomized trials of palliative RT in 114 NSCLC patients showed no spinal myelopathy when treated with 10Gy in one fraction. However patients treated with 17Gy in 2 fractions had a cumulative risk of 2.2 % of developing myelopathy at 2 years. Further data has shown that Spinal cord can tolerate 10Gy to 10% of the volume as defined as 6 mm above and below the target lesion with acceptable rates of myelitis [13].

A study by Collins et al. [14] treated 20 patients with inoperable Stage 1 NSCLC with CK in Georgetown University Hospital, Washington DC, US. Dose ranged from 42-60Gy in 3 fractions and median follow up of surviving patients was 25 months with an overall survival estimate of 87%. The present study also reported doses for primary and metastatic lung cancers range from 54 - 60Gy in 3#.

Spinal tumours/metastases and Re-irradiation studies:

A number of studies have shown no radiation induced myelopathy after a Biological effective dose of 80-100Gy to spinal cord at a median follow up of 8 months [15-17]. Patients receiving BED > 102 Gy seems to show myelopathy [18]. Another study has concluded that a point maximum dose of 10Gy is safe as radiation induce myelopathy was found to take place when maximum point doses are 14.8, 13.1 and 10.6 Gy in a single fraction [19].

In the present study either no margin or a margin of 1mm was used around CTV for treatment of spinal metastases with a dose of 27Gy in 3 # (9 Gy /#). This seems to be safe dose with probably a low and acceptable cumulative risk of myelopathy, with high probability of tumour control and symptom relieve.

Yamada et al.[20] reported no myelopathy or other late toxicities in 93 patients that were treated with a median dose of 24 Gy (range 18-24Gy) with spinal cord maximum point dose restricted to 14 Gy. After a median follow up of 15 months, the actuarial 1 year control rate was 90%. This study found a direct dose response relationship i.e. higher doses give rise to better local control rates. The spinal radiosurgery was conducted in Memorial Sloan-Kettering hospital.

A phase I/II trial conducted at the MD Anderson Cancer centre treated 63 patients with hypofractionated course of spinal radiosurgery with a fractionated regime of 6 Gy in 5 fractions to half the patients and 9 Gy in 3 fractions given to other half. No grade 3 or 4 neurologic toxicity or myelopathy was reported with a median follow up of 21 months and the one year actuarial progression-free rate was 84%. The study reported one case of grade 3 nausea, vomiting and diarrhea, one case of grade 3 dysphagia and trismus and one case of grade 3 non cardiac chest pain. The study recommended using wide posterior margin to diseased vertebrae to avoid recurrence in bone adjacent to the spinal cord and in epidural space [21].

Brain tumour studies:

A study by yang et al. [22] showed that CyberKnife treatment is effective in treatment of metastatic brain disease. A patient with more than 24 brain lesions was treated with CyberKnife and was given a total dose of 22Gy in 3 fractions showed complete disappearance of the tumour 3 months post treatment

A retrospective study by Acker et al. [23] showed safety and efficacy of CyberKnife treatment in elderly patients with brain metastases. The projected overall survival at 3, 6 and 12 months after treatment were 79, 55 and 23% respectively while the and local tumour progression free survival at 6, 12, 36 and 72 months post treatment were 99.2, 89.0 and 67.2, 64.6 and 64.6% respectively. The predictive factors for local progression were Older age and female sex. The study reported Karnofsky performance score remained steady in 97.9% of the patients.

Another study by Telentschak et al. [24] reported actuarial local control rates at 3, 6, and 12 months were 98%, 98%, and 78.6%, respectively in patients with critical brain metastases. 12 % of patients had grade I to III complications. The study found that median overall survival was associated with higher KPS.

Liver tumours studies: A study conducted by Kato et al. [10] 65 advanced and terminal stage HCC patients (with 95 lesions) with CyberKnife and reported better survival with doses greater than or equal to 30 Gy. Out of 52 cases of bone metastases, 69% of patients achieved pain relief. Toxicity included grade 4 Cerebral bleeding in one patient treated for brain metastases and grade 2 oesophageal ulcer in another patient post treatment who was treated for hepatic vessel lesion (Complete response was achieved with 31.2Gy to oesophagus) The Treatment Planning system (TPS) used was MultiPlan[®] (Accuray) and Synchrony[®] (Accuray) tracking system was used to track the tumour. The Planning target volume for intra-hepatic lesions and lung metastases include GTV plus 2-5 mm margin in all directions whereas the PTV for spinal lesions included GTV plus 2 mm margin and for brain metastases no margins were applied to GTV. Total dose ranged from 8-50Gy, delivered in 1-10 fractions and prescribed to the 80% isodose line administered to the PTV over 1-7 consecutive working days. The median prescribed dose for tumours invading hepatic vessels or bile duct was 35 (range: 28-50 Gy) in 3-10 fractions where as median prescribed dose for extrahepatic lesions was 25 Gy (6-48) in 1-6 fractions. The response rate was 48% and disease control rate was 76% for all lesions after excluding unevaluated cases. The response rate and disease control rates for tumours invading the hepatic vessels or bile duct were 50% and 80% respectively. As far as adverse effects are concerned no patient had a grade 2 or higher toxicity. No classic Radiation induced Liver disease, considerable rises in liver enzyme and haemotologic complications were detected during treatment. Compared with these results, the margins and doses reported in the present case studies are similar. The most common dose was 45Gy in 3 fractions for primary liver tumours and 54Gy in 3 fractions for liver metastases in the present study which is above 30Gy. However doses for spinal metastasis were 27Gy in 3 fractions which is more than the median dose used for extrahepatic lesions in Kato et al study but lower than 30Gy.

A study **by Kang et al. [25] o**bserved Response rate of 66.7% for portal vein tumour thrombosis treated by SBRT alone and authors suggested that response rates of up to 73.5% could be achieved if combined with TACE (trans arterial chemo-embolization).

Another study conducted **by Goyal et al. [26]** involving unresectable liver tumours reported a 60% mean decrease in tumour volume three months post-treatment in case of HCC patients whereas a mean reduction in tumour volume of 59% was observed three months post treatment in case of liver metastases. Initial control rate was 82% with a median follow up of 8 months and three patients (two liver metastases patients and one patient with IHC) suffered from recurrences while seven patients experienced distant recurrences. The median **prescribed dose was 34Gy** (24-45Gy) in 1-3 fractions prescribed to median prescription isodose line of 70%. The study reported two grade 2 Gastrointestinal ulcers and one grade 3 GI ulcer. The authors concluded that CyberKnife Stereotactic Radiosurgery is successful local treatment for unresectable tumours of the liver.

Pancreatic tumour studies:

A study by song et al. [9] assessed the efficacy and safety of CyberKnife treatment for locally advanced pancreatic tumours (LAPC) and reported the median OS of 12.5 months and 1 year and 2 year survival rates of 53.9% and 35.1% respectively with one year freedom from local progression (FFLP) rate of 90.8% when treated with a **median dose of 45Gy (35Gy – 50Gy)** in 5 fractions. 61% of the patients experienced Grade 1-2 acute and late stage GI reactions where as one patient suffered from grade 3 toxicity. Multiplan Treatment planning system was used to create CyberKnife treatment plans and PTV was obtained by **adding a 3mm margin to GTV**. The **CK Synchrony motion tracking system** was used along with fiducial markers. The margins and dose reported in the song et al study are similar to the present case studies. Both respondents from US and Australia reported using 3-5 mm margin around GTV to obtain PTV for localized pancreatic cancer. Dose used in Australian institute was 40Gy in 5 fractions.

A study by Ji et al. [27] that compared CK SBRT plus Chemotherapy with Chemotherapy alone found that addition of SBRT improved local control rate (6 month PFS rate was 29.4% vs. 20.6% in CK+Chemotherapy and chemotherapy group alone)) but did not improve overall survival in patients with primary tumour of Liver only oligometastatic pancreatic cancer, primarily because many patients suffered from distant metastasis. There was no significant difference in the toxicity between the two groups.

PC studies:

A systematic review that assessed the clinical evidence of gantry versus robotic arm SBRT in prostate cancer patients concluded that neither device could be advocated for all prostate cancer patients [28]. However Robotic SBRT resulted in better or comparable freedom from biochemical failure for low and intermediate risk prostate cancer patients at 5-7 years. In terms of acute and late toxicities Robotic SBRT and Gantry based SBRT showed comparable results. The gantry based treatment resulted in grade 2 and greater GU toxicities that ranged from 5-8% vs. 4% -19.2% toxicity with Robotic SBRT. The GI grade 2 and greater toxicities in gantry based studies ranged from 7.5% - 8% vs. 0-12% in Robotic SBRT studies. while interpreting these results it is important to note that gantry based studies only had low risk patients and only 3 studies were reviewed whereas Robotic based studies included low, intermediate and high risk patients. The longer follow up and more extensive quality of life studies might change the reported toxicity percentages. The authors concluded that gantry based SBRT could be more useful for obese patients as higher energies could be used to treat these patients (greater than 6MV) and gantry based SBRT offers shorter treatment time per fraction compared to robotic SBRT [28]. The dose ranged from 33.5Gy to 40 Gy in 5 fractions in Gantry based studies and 32Gy-40Gy in 4-5 fractions in robotic SBRT studies.

Another study compared the CK plans with IMRT based techniques (VMAT, IMRT Sliding window, Helical Tomotherapy) for prostate cancer patients [29]. The study found no dosimetric differences in terms of PTV coverage and conformality but better PTV homogeneity was observed with rotational IMRT techniques at medium and high dose range. Bladder and rectum sparing was again better achieved with IMRT techniques than CK [29]. Helical Tomotherapy showed superior Normal Tissue Complication Probablity (NTCP) for rectum but no difference was observed for NTCP values for bladder with any of the techniques. The target dose used in this study was 36.25Gy in 5 fractions over 1 week which is the same dose reported by professionals in the present study. As far as margins are concerned the present study results are in agreement with the margins used by Scobioala et al. [29]. In Scobioala et al. [29] study CTV included only Prostate (no Seminal vesicles) and to obtain PTV a 3mm margin was added in the dorsal direction (posteriorly) and a 5mm margin in ventral (Anteriorly) and lateral directions.

Some researchers thinks CK may be associated with higher secondary malignancy rates due to a large volume of normal tissue receiving low dose radiotherapy along with longer treatment times and higher Monitor Units given by CK [30]. Researchers have suggested algorithms that can be used to reduce treatment delivery time by using beam angle class solutions for non coplanar SBRT with CK rather than using beam angle optimization for each individual patient [31].

A study by Rossi et al also showed superiority of Automatically generated CK robotic plans over manually generated CK plans. AutoROBOT CK plans produced better rectal sparing than automatically generated VMAT plans [32].

40 | Page

Reasons behind CK adoption:

As far as reasons of CK adoption are concerned both experts agreed that CK was adopted to provide precise treatment delivery, to achieve better local control rates, to give re-treatment and to gain competitive edge in the clinical practice. The CK expert from Australia provided additional reasons for CK adoption namely dose escalation, reduce treatment time and for clinical research purposes. A study by Brown et al.[6] showed that all NSCLC patients except one achieved at least partial response (30% reduction in tumour) and concluded that excellent control rates were achieved in early stage NSCLC patients when treated with CyberKnife.

A retrospective study conducted by Liu et al. [33] to evaluate safety and efficacy of CK treatment in 13 patients with olfactory groove meningiomas found 12 out of thirteen patients achieved 100% regional control rate at the time of follow up. There was a median tumour volume reduction of 31.7%. The study employed three dose regimes depending on tumour size i.e. 10Gy in 1fraction for tumours less than 10 cm³, 25Gy in 5 fractions and 54Gy in 30 fractions for tumours greater than 10 cm³ or in close vicinity of OARs. This study was conducted a medical centre in Boston, USA.

A study by Jereczek-Fossa et al. [8] found Actuarial 3 year in field progression free survival of 67.6%, Progression free survival of 18.4% and Overall survival of 31.2% in oligometastatic cancer patients treated with CK. The median dose was 24Gy in 3 fractions and complete radiological response was recorded in 17% of the lesions and partial response in 29% of the lesions. In 39% of the lesions the disease was found to be stabilized while in 15% of the lesions progressive disease was observed. The study concluded that CK treatment gives long term in-field tumour control with low toxicity.

A study by song et al. [9] showed median overall survival of 12.5 months in patients with locally advanced pancreatic cancer. 53.9 % of the patients had OS of one year whereas 35.1% of patients has a 2 year OS. The study reported 1 year freedom from local progression of 90.8%. This study treated patients with a median dose of 45Gy in 5 fractions whereas the prescribed dose ranged from 35-50Gy in 3-8 fractions. 90% of patients received Chemotherapy before or after CK treatment and grade 1-2 acute and late Gastrointestinal toxicity was reported in 61% of patients. In the present case study the most common dose regime used for pancreatic cancers was 40Gy in 5 fractions in Australian CK centre which is in line with the study by Song et al. [9].

A Case study conducted by **Accuray in St.Joseph's Hospital**, Phoneix, Arizon, US observed radiographically complete response in a patient suffering from $T1N_0\ M_0\ NSCLC$ three months post treatment [34]. The patient had no surgery and was treated with 48Gy in 3 fractions (16Gy/fraction) while tumour motion was managed by CK Synchrony tracking system. A 5mm margin was added to GTV to get PTV. This study involved a radiation oncologist, a medical physicist and a Radiation therapist. This is in line with the present study as both respondents from US and Australia reported using a 5mm Margin to get PTV while planning CK treatment for lung cancer patients.

Another case study conducted by Accuray [35] in CyberKnife center of Miami, USA showed no evidence of disease 11 months post treatment with Fiducial free CK for T1N0M0 NSCLC. The patient was treated with 60Gy in 3 fractions (20Gy/Fraction) and motion was managed by XSight Lung tumour tracking system (Synchrony). The CK team in this case study included one radiation oncologist, one Thoracis Surgeon, two physicists, one dosimetrist and 2 therapists. The dose in the accuracy case study is similar to the dose usage reported by US CK expert in the present study for treatment of early stage Lung cancer.

CK Team composition

The data regarding number and type of CK team is mixed. However it seems that CK team must include at minimum a Medical Physicist, 2 Radiotherapy technologists and 1 radiation oncologist.

Tumour Tracking system

The present study has also showed that Synchrony system of CyberKnife is used in lung, pancreas, liver and kidney cancer patients for motion management in both Australia and USA. This is in agreement with the literature. A study by Nuyttens and Pol [36] showed CK synchrony system (4D rea time tumour tracking) can be used to treat moving tumours with 2mm accuracy while patients breathe normally.

CK Training Pathways

The present study has shown that in Australia SBRT symposium is used to gain CK knowledge and experience where as in USA the emphasis is on manufacturer's training programmes. No studies could be found that describe what strategies are used to gain CK Knowledge and experience in clinical and industry setting. The expert from the Australia also mentioned Experience as one of the ways to gain CK experience. Author of the present study assumes that he meant probably in house training. Author of the current study recommend using other strategies to improve CK knowledge and experience of staff and radiation oncology students such as by offering Mentor based training, by designing and offering university courses that meet industry needs, by

offering practical hand on experiences in workshops, by encouraging oncology, medical physics and radiography related societies (e.g. ASCO, ESTRO, RTOG, APS, AAPM) to offer clinically relevant courses and workshops, by offering internships in Medical physics and by including physics and dosimetry in Undergraduate and post graduate syllabuses.

Future Directions

For future studies, author recommends doing similar studies but involving multiple institutes in USA, Australia, Europe and Asia to make data more generalizable and to gain more information on treatment induced toxicity, Local failure rates, overall survival, CK related organizational resources as well as on quality of life of cancer patients who have undergone CyberKnifetreatment

V. CONCLUSION

The present study consists of two comparative case studies and provides an overview of clinical, technical, organizational and Educational strategies and resources used by two institutes in USA and Australia to provide Stereotactic Radiosurgery and Stereotactic body Radiotherapy to cancer patients. The study captures the perspectives of two CyberKnife experts who have considerable experience of using CyberKnife.

As far as clinical resources are concerned the data regarding number and type of CyberKnife team is mixed. However it seems that CK team must include at minimum one Medical Physicist, two Radiotherapy technologists and 1 radiation oncologist. With respect to clinical treatment intent, CK is used to provide curative treatment in American institute and curative as well as palliative treatment in Australian institute. The study has shown CK is used for both intra and extra cranial tumours in both institutes in USA and Australia. In USA CK is used for only early stage disease whereas it is used for both early and advanced stage cancers in Australian institute.

In terms of **technical resources**, CyberKnife Multiplan Treatment planning system is used by Medical physicists to create CK treatment plan in US institute whereas Accuray Precision Radiotherapy TPS are used by Radiation technologists to create CK treatment plans in Australian institute. The study has shown that In room volumetric imaging, CyberKnife tracking system (6D Skull, Fiducial, X Sight spine with Synchrony, X Sight lung with Synchrony and Fiducials) and planar imaging are used before and during the treatment to localize and verify the target based on various sites. In US institute, Fiducial markers are not used for brain, spine and most lung tumours. In terms of immobilization devices mask for c-spine and brain tumours are used in US institute and headframe are used during treatment of brain tumours in Australian institute.

In the present study Different doses were reported for lung cancers by both experts (60 Gy in 3-5# in USA vs. 54Gy in 3# Australia). Most common dose was spinal cord tumours was 27Gy in 3 fractions, for liver tumours 45 Gy - 54 Gy in 3 fractions, for prostate cancer was 35-36Gy in 5 fractions and for pancreatic tumours 40Gy in 5 fractions. In the present study fatigue and tiredness were most common acute toxicity experienced by patients suffering from prostate, liver, pancreas and kidney cancers.

In the present study zero margin was added to GTV to obtain CTV for localized prostate disease and a margin of 3mm posteriorly and 5mm anteriorly and laterally was added to obtain PTV. Both respondents from US and Australia reported using 3-5 mm margin around GTV to obtain PTV for localized pancreatic cancer. In the present case studies a margin of 2mm was added to achieve PTV for spinal metastases in USA case study and a margin of zero in Australian case study. A margin of 5mm was reported in both case studies for liver metastasis.

Under the category of CK organizational resources and strategies, the present study found challenges faced during CK implementation in the organization and reasons for CK adoption. As far as reasons of CK adoption are concerned both experts agreed that CK was adopted to provide precise treatment delivery, to achieve better local control rates, to give re-treatment and to gain competitive edge in the clinical practice. The CK expert from Australia provided additional reasons for CK adoption namely dose escalation, reduce treatment time and for clinical research purposes. In the present study, the respondent from Australia said they experienced financial difficulties while respondent from US said they experienced no challenges in implementing CK in the institute. Both CK experts found CK technology cost effective compared to VMAT, IMRT and Tomotherapy. From the perspectives of US CK expert, CK provides better tumour control, DFS, OS and reduced treatment induced toxicities compared to IMRT,VMAT and proton therapy.

To ensure accurate and efficient CK implementation, treatment planning, delivery and Quality assurance staff must be well educated. The present study has shown that in Australia SBRT symposium is used to gain CK knowledge and experience where as in USA the emphasis is on manufacturer's training programmes. The study also found that In USA Radiation oncology and medical physics programmes provide SBRT and CK experience whereas according to CK expert in Australian institute oncology and medical physics programmes do not provide SBRT and CK experience.

Author of the current study recommend using other strategies to improve CK knowledge and experience of staff and radiation oncology students such as by offering Mentor based training, by designing and offering university courses that meet industry needs, by offering practical hand on experiences in workshops, by encouraging oncology, medical physics and radiography related societies (e.g. ASCO, ESTRO, RTOG, APS, AAPM) to offer clinically relevant courses and workshops, by offering internships in Medical physics and by including physics and dosimetry in Undergraduate and post graduate syllabuses. In summary, this study shows similarities and dissimilarities involving the use of CK technology in two institutes in USA and Australia.

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Conflict of Interest

Author discloses no conflict of Interest.

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Supplementary Data: Supplementary data related to this article can be found below:

Appendix A: A sample CyberKnife E- Survey 2020

| Demogr | aphic Information | Q.6. How | many professionals are involved in CK treatment planning and delivery (e.g. Radiation oncologist : 1)? |
|--|---|------------|--|
| | | Α. | Medical physicists: |
| Q.1 Gen | der of the Respondent | В. | Radiographers: |
| | | C. | Radiation technologists: |
| Α. | Male | D. | Dosimetrists: |
| В. | Female | E. | Radiation oncologists: |
| Q.2 Age | | Q. 7. Loca | ation: city, country of cyber knife (CK) institution? |
| A. | Less than 30 years 30-40 years | | |
| B. C. | 40-50 years | Q.8. What | challenges did you or your institution faced when implementing cyber knife? Please select all options that apply to you. |
| D. | 50-60 years | | |
| E. | Above 70 years | A. | Lack of knowledge/ Difficulty acquiring required CK knowledge |
| | tal status | B. | Lack of training |
| Α. | Single | C. | Shortage of guidelines for cyber knife treatment planning for various cancers |
| В. | Married | D. | Shortage of QA guidelines for cyber knife setup and implementation. |
| C. | Divorced | E. | Financial |
| D. | 4. Separated | F. | Lack of personnel |
| Q.4 Occ | upation (Please state your occupation): | | |
| A. | Radiation oncologist | G. | Complex QA/Commissioning programme |
| B. | Medical Physicist | Η. | Others (please specify): |
| C. | Radiation/ Radiotherapy Dosimetrist | Informatic | on about Cyber knife |
| D. | Research medical physicist | milorinaci | and about dybet name |
| E | Director | 000 | ou have cyber knife and Stereotactic RT in your hospital or institution? |
| F. | Other (Please specify): | Q.5. D0 y | ou have cycler knite and Stereotactic KT in your hospital or institution? |
| Cyber kı | nife institutional background | A. | Yes |
| | In a | B. | No. |
| Committee of the Commit | of Practice | C. | |
| A. | Academic practice/ centre | C. | Do not know |
| В. | Private practice/ centre | O 10 Have | v long you have been using cyber knife and SBRT to give treatment? |
| C. | Public | Q.10. HO | violig you have been using cyber knille and obtain to give treatment? |
| D. | Not for profit | A | Less than 6 months |
| E. | Other: please specify | | |
| F. | Do not know | В. | 1 year |
| | PSO(00004-0200). | C. | 5 years |
| | | D. | 10 years |
| | | E. | Others (please specify) |
| | | 044.5 | |

| A. Intracranial B. Body tumo C. Intracranial D. Do not kno | urs i and body tumours w | ife? Do you ! | reat Primary or m | etastatic lesions? | (e.g. primary NSCLC, liver n | netastases, spinal | | Q.15. | Do you use cyber k A. Early stage (ES B. Advanced stag C. Do not know Why did you adopt! A. To achieve dos B. To achieve mo C. To get better lo D. To reduce treat | c (AS) Cyber knife? e escalation re precise treatment cal control rates | | r advanced sta | age disease? |
|---|--|------------------|------------------------|----------------------|------------------------------------|--------------------|---|--------------------------|--|--|-------------------------|---------------------------------|------------------------------|
| | | Primary | lesion/tumour | | Metastases | | | | E. For retreatmen F. Clinical research | h | | | |
| Ca Lung | | | | | | | | | G. To gain a compH. Others (Please | etitive edge | | | |
| Ca Prostate (PC) | | | | | | | | | ment Planning and D | | | | |
| Ca liver (HCC) | | | | | | | | Q. 16 | What CK image gu | idance system o | onsists of? | | |
| Ca pancreas | | | | | | | | | | | | | |
| Spinal cancer | | | | | | | 147 Do um um | a section of in | amabilization desire where | union CobarKolle t | continuent editamina su | nd delivery (e.a. | SBF = stereotactic body fram |
| Brain cancer | | | | | | ě | tc)? If yes whic | th one? | | | | | |
| Kidney cancer | | | | | | | Calung | | Head Frame | SBF | Alpha cr | radle | Other (Please specify) |
| Others (Please speci | Mi- | | | | | | Ca Prostate (P | c) | | | | | |
| Others (r lease specif | 97/- | | | | | | Ca liver (HCC) | | | | | | |
| | | | | | | | Ca pancreas | | | | | | |
| | | | | | | | Spinal cancer | | | | | | |
| | | | | | | | Brain cancer | | | | | | |
| | | | | | | | Kidney cancer Others (Please | specifyl | | | | | |
| | | | | | | | | | | | | | |
| CK image Fiducial n SBF to es | om volumetric and plan e guidance system narkers stablish an external co o ease specify. | | n. | | | Liver me | ctable HOC estastases ent: Unresect | able | | | | | |
| Do not kn | | | | ev. | | Q.22. Wh | sat is the mos | t common sch 16Gy ir | edule used for primary a | nd metastatic spin. 18Gy in 18 | al lesions when tre | oating with CK? Other (Pleas | se specify) |
| Q. 19. What is the most co | enmon schedule used for P | rostate cancer (| PC) when treating with | CK | | | spinel lesions | | | | | | |
| | 35-36 in 5# | 46Gy in 5# | 47.5Gy in 5# | 50Gy in 5# | Other (Please specify) | Metasta | atic spinal lesio | ns | | | | | |
| Primary localized PC (71-72) | | | | | | | | | | | | | |
| Metastatic PC | | | | | | Q. 23. W | hat is the mo | est common s | chedule used for prima | ary and metastatic | pancreatic lesion | ns when treating | g with CK? |
| | | | | | | 0 | | and a | | | | | |
| Q. 20. What is the most co Advanced stage (Stage III | ommon schedule used for p I-IV). | rimary and met | astatic lung lesion wh | en treating with CK? | ES= Early stage (Stage I-II), AS= | Primary lesions | y panci | esec | | | | | |
| | 35-36 in 5# | 46Gy in 5# | 47.5Gy in 5# | 50Gy in 5# | Other (Please specify) | Metasta lesions | | readic | | | | | |
| Primary lung tumour, ES | | | | | | | | | | | | | |
| Primary lung lumour, AS | | | | | | | | | | | | | |
| Metastatic lung cancer | | | | | | | | | | | | | |
| Recurrent lung cancer | | | | | | | | | | | | | |
| Q. 24. How much ma | rgin you apply to GTV | to get CTV a | and PTV in each of | the following ca | ses? ES= Early stage, | | Q. 2 | 7. Who carr A. Med | ies out CK radiation ical Physicists | treatment plans | ning in your org | panization? | |
| AD= Advanced stage | e, HCC= Hepatocellula | r carcinoma, | PC= Prostate can | cer. | | | | B. Rad | ographers ation technologists | | | | |
| | GTV | (| TV = GTV+ margi | n PTV= CT | V+ margin | | | D. Rad | ation technologists ation Dosimetrists ics Assistant | | | | |
| Primary localized (71-72) | PC (| | | | | | 111220 | | | | | | |
| | | | | | | | Q.28 | . Which typ | e of techniques you Ca Lung | employ to cont Ca PC | | nduced motion tas Ca live | |
| Metastatic PC | | | | | | | Fie | fucial marke. | | | 8) | | 70 |
| Primary ES Ca lung | | | | | | | | spiratory ga | | | | | |
| Primary AD ca lung | | | | | | | | | wy | | | | |
| Primary small HCC | | | | | | | | dominal mpression | | | | | |
| Primary large HCC | | | | | | | Re | al time t | umour | | | | |
| Metastatic HCC | | | | | | | | cking | | | | | |
| Spinal metastases | | | | | | | | | | | | | |
| Liver Metastases | | | | | | | CA | | ystem | | | | |
| Liver metastases | | | | | | | (S) Re | ynchrony spiratory tr | acking | | | | |

2.26. Which treatment planning systems you use to create radiation plans for Cyber knife?

- A. Multiplan treatment planning systems you
- C. Edipsi
- D. Other (Please specify

Q. 29 What most common acute toxicities patients experience (up to 3 months) after receiving CK treatment in each case:

| | Ca lung | Ca PC | Ca liver | Ca Pancreas | Ca kidney | Spinal ca |
|----------------------|---------|-------|----------|-------------|-----------|-----------|
| Dyspnea | | | | | | |
| Pneumonitis | | | | | | |
| Diamhea | | | | | | |
| Other Bowel problems | | | | | | |
| Bladder problems | | | | | | |
| atique & Tiredness | | | | | | |
| Pain | | | | | | |
| Appetite loss | | | | | | |

Knowledge and Experience:

- Q.30. How did you and other team members in your organization gain the required CK and SBRT knowledge and experience?
 - A. Manufacturer training programme
 - B. SBRT symposium
 - C. University courses
 - D. Other: (Please specify)
- Q. 31. Are radiation oncology and medical physics programs in your country provide experience in SBRT and CK?
- Q.32. Is CK treatment cost effective compared to other Radiation therapy technologies such as VMAT, IMRT, Tomotherapy?
- Q.33 For which cancers CK treatment provides better tumour control, disease free survival and overall survival and reduced treatment induced morbidity compared to IMRT, VMAT and proton therapy?

REFERENCES

- [1]. Accuray. CyberKnife® S7TM Precise Robotic Radiotherapy treatment as individual as every patient [Online]. 2021. Available at: https://www.accuray.com/cyberknife/ [Accessed 16/10/2021].
- [2]. Guckenberger M, Andratschke N, Alheit H, Holy R, Moustakis C, Nestle U, Sauer O. Definition of stereotactic body radiotherapy. Strahlentherapie und Onkologie. 2014;190(1):26-33. [online] Available at: https://link.springer.com/content/pdf/10.1007/s00066-013-0450-y.pdf [Accessed 23/10/2021]
- [3]. Kurup G. CyberKnife: A new paradigm in radiotherapy. *Journal of medical physics/Association of Medical Physicists of India*. 2010;35(2):63. [online] Available at: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2884306/ [Accessed 23/10/2021]
- [4]. Eken S, Zorlu F, Yeginer M, Ozyigit G. Performance evaluation of the X-sight spine tracking system for abdominal tumors distal to spine: A 2D dosimetric analysis. *Medical Dosimetry*. 2019;44(4):370-4.
- [5]. Dieterich S. and Pawlicki T. (2008) CyberKnife Image Guided Delivery and Quality Assurance, *International Journal of Radiation Oncology Biology Physics*, 71 (1), S126-130 [online] Available at: https://www.sciencedirect.com/science/article/pii/S0360301607042939 [Accessed 21/09/2021
- [6]. Brown W, Wu X, Wen B, Fowler J, Fayad F, Amendola B, García S, Zerda AD, Huang Z, Schwade J. Early results of CyberKnife image-guided robotic stereotactic radiosurgery for treatment of lung tumors. Computer Aided Surgery. 2007;12(5):253-61. [online] Available at: https://www.tandfonline.com/doi/full/10.3109/10929080701684754 [Accessed 23/10/2021]
- [7]. Conti A, Pontoriero A, Arpa D, Siragusa C, Tomasello C, Romanelli P, Cardali S, Granata F, De Renzis C, Tomasello F. Efficacy and toxicity of CyberKnife re-irradiation and "dose dense" temozolomide for recurrent gliomas. *Acta neurochirurgica*. 2012:154(2):203-9.
- [8]. Jereczek-Fossa B, Bossi-Zanetti I, Mauro R, Beltramo G, Fariselli L, Bianchi L, Fodor C, Fossati P, Baroni G, Orecchia R. CyberKnife robotic image-guided stereotactic radiotherapy for oligometastic cancer. Strahlentherapie und Onkologie. 2013;189(6):448-55.
- [9]. Song Y, Yuan Z, Li F, Dong Y, Zhuang H, Wang J, Chen H, Wang P. Analysis of clinical efficacy of CyberKnife® treatment for locally advanced pancreatic cancer. *OncoTargets and therapy*. 2015;8:1427.
- [10]. Kato H, Yoshida H, Taniguch H, Nomura R, Sato K, Suzuki I, Nakata R. CyberKnife treatment for advanced or terminal stage hepatocellular carcinoma. *World journal of gastroenterology*. 2015;21(46):13101.
- [11]. Rades D, Stalpers L, Veninga T, Schulte R, Hoskin P, Obralic N, Bajrovic A, Rudat V, Schwarz R, Hulshof M, Poortmans P. Evaluation of five radiation schedules and prognostic factors for metastatic spinal cord compression. *Journal of Clinical Oncology*. 2005;23(15):3366-75.
- [12]. Rades D, Stalpers L, Hulshof M, Zschenker O, Alberti W, Koning C. Effectiveness and toxicity of single-fraction radiotherapy with 1×8 Gy for metastatic spinal cord compression. *Radiotherapy and oncology.* 2005;75(1):70-3.
- [13]. Ryu S, Jin J, Jin R, Rock J, Ajlouni M, Movsas B, Rosenblum M, Kim J. Partial volume tolerance of the spinal cord and complications of single-dose radiosurgery. Cancer: *Interdisciplinary International Journal of the American Cancer Society*. 2007;109(3):628-36.

- [14]. Collins B, Vahdat S, Erickson K, Collins S, Suy S, Yu X, Zhang Y, Subramaniam D, Reichner C, Sarikaya I, Esposito G. Radical CyberKnife radiosurgery with tumor tracking: an effective treatment for inoperable small peripheral stage I non-small cell lung cancer. *Journal of hematology & oncology*. 2009;2(1):1-9.
- [15]. Brenner D. The linear-quadratic model is an appropriate methodology for determining isoeffective doses at large doses per fraction. Seminars in radiation oncology. 2008;18(4):234-239.
- [16]. Jones L, Hoban P, Metcalfe P. The use of the linear quadratic model in radiotherapy: a review. Australasian Physics & Engineering Sciences in Medicine. 2001;24(3):132-46.
- [17]. Rades D, Stalpers L, Veninga T, Hoskin P. Spinal reirradiation after short-course RT for metastatic spinal cord compression. International Journal of Radiation Oncology* Biology* Physics. 2005;63(3):872-5.
- [18]. Nieder C, Grosu A, Andratschke N, Molls M. Proposal of human spinal cord reirradiation dose based on collection of data from 40 patients. *International Journal of Radiation Oncology* Biology* Physics*. 2005 Mar 1;61(3):851-5.
- [19]. Sahgal A, Ma L, Gibbs I, Gerszten P, Ryu S, Soltys S, Weinberg V, Wong S, Chang E, Fowler J, Larson D. Spinal cord tolerance for stereotactic body radiotherapy. *International Journal of Radiation Oncology* Biology* Physics*. 2010;77(2):548-53.
- [20]. Yamada Y, Bilsky M, Lovelock D, Venkatraman E, Toner S, Johnson J, Zatcky J, Zelefsky M, Fuks Z. High-dose, single-fraction image-guided intensity-modulated radiotherapy for metastatic spinal lesions. *International Journal of Radiation Oncology* Biology* Physics*. 2008;71(2):484-90.
- [21]. Chang E, Shiu A, Mendel E, Mathews L, Mahajan A, Allen P, Weinberg J, Brown B, Wang X, Woo S, Cleeland C. Phase I/II study of stereotactic body radiotherapy for spinal metastasis and its pattern of failure. *Journal of Neurosurgery*. 2007;7(2):151-60.
- [22]. Yang G, Wang Y, Wang Y, Lin S, Sun D. CyberKnife therapy of 24 multiple brain metastases from lung cancer: A case report. Oncology letters. 2013;6(2):534-6.
- [23]. Acker G, Hashemi SM, Fuellhase J, Kluge A, Conti A, Kufeld M, Kreimeier A, Loebel F, Kord M, Sladek D, Stromberger C. Efficacy and safety of CyberKnife radiosurgery in elderly patients with brain metastases: a retrospective clinical evaluation. Radiation Oncology. 2020;15(1):1-0. [online] Available at:https://ro-journal.biomedcentral.com/articles/10.1186/s13014-020-01655-8 [Accessed 21/10/2021]
- [24]. Telentschak S, Ruess D, Grau S, Goldbrunner R, von Spreckelsen N, Jablonska K, Treuer H, Kocher M, Ruge M. Cyberknife® hypofractionated stereotactic radiosurgery (CK-hSRS) as salvage treatment for brain metastases. *Journal of Cancer Research and Clinical Oncology*. 2021;147:2765-2773. [online] Available at: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8310836/ [Accessed 21/10/2021]
- [25]. Kang J, Nie Q, Du R, Zhang L, Zhang J, Li Q, Li J, Qi W. Stereotactic body radiotherapy combined with transarterial chemoembolization for hepatocellular carcinoma with portal vein tumor thrombosis. *Molecular and clinical oncology*. 2014;2(1):43-50.
- [26]. Goyal K, Einstein D, Yao M, Kunos C, Barton F, Singh D, Siegel C, Stulberg J, Sanabria J. CyberKnife stereotactic body radiation therapy for nonresectable tumors of the liver: preliminary results. HPB Surgery. 2010;2010: 1-8 [online] Available at: https://downloads.hindawi.com/archive/2010/309780.pdf [Accessed 23/10/2021]
- [27]. Ji X, Zhao Y, He C, Han S, Zhu X, Shen Z, Chen C, Chu X. Clinical effects of stereotactic body radiation therapy targeting the primary tumor of liver-only oligometastatic pancreatic cancer. Frontiers in oncology. 2021;11;659987. [online] Available at: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8190391/ [Accessed 23/10/2021]
- [28]. Avkshtol V, Dong Y, Hayes S, Hallman M, Price R, Sobczak M, Horwitz E, Zaorsky N. A comparison of robotic arm versus gantry linear accelerator stereotactic body radiation therapy for prostate cancer. *Research and reports in urology.* 2016;8:145-158. [online] Available at: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4993397/ [Accessed 09/10/2021].
- [29]. Scobioala S, Kittel C, Elsayad K, Kroeger K, Oertel M, Samhouri L, Haverkamp U, Eich H. A treatment planning study comparing IMRT techniques and cyber knife for stereotactic body radiotherapy of low-risk prostate carcinoma. *Radiation Oncology*. 2019;14(1):1-0. [online] Available at: https://ro-journal.biomedcentral.com/articles/10.1186/s13014-019-1353-6 [Accessed 09/10/2021].
- [30]. Lin Y, Lin K, Ho H, Lin L, Lee S, Chui C. Treatment plan comparison between stereotactic body radiation therapy techniques for prostate cancer: non-isocentric CyberKnife versus isocentric RapidArc. *Physica Medica*. 2014;30(6):654-61.
- [31]. Rossi L, Breedveld S, Aluwini S, Heijmen B. Noncoplanar beam angle class solutions to replace time-consuming patient-specific beam angle optimization in robotic prostate stereotactic body radiation therapy. *International Journal of Radiation Oncology* Biology* Physics*. 2015;92(4):762-70.
- [32]. Rossi L, Sharfo A, Aluwini S, Dirkx M, Breedveld S, Heijmen B. First fully automated planning solution for robotic radiosurgery–comparison with automatically planned volumetric arc therapy for prostate cancer. *Acta Oncologica*. 2018;57(11):1490-8.
- [33]. Liu J, Rojas R, Lam F, Mirza FA, Mahadevan A, Kasper E. Indications, feasibility, safety, and efficacy of CyberKnife radiotherapy for the treatment of olfactory groove meningiomas: a single institutional retrospective series. Radiation Oncology. 2020;15(1):1-0.
- [34]. Accuray Incorporated. Case Study. Non Small Cell Lung Cancer. Left Upper Lung. [online].2007.Available at: https://cyberknife.com/wp-content/uploads/500093-A-NonSmallCell-Lung-Cancer.pdf [Accessed 23/09/20221]
- [35]. Accuracy Incorporated. CaseStudy. Non invasive Stereotactic Radiosurgical Treatment of Non Small cell lung cancer. [online]. n.d. Available at: https://www.accuray.com/wp-content/uploads/500341.b_xsight_case_study.pdf [Accessed 23/09/2021]
- [36]. Nuyttens J, Van De Pol M. The CyberKnife radiosurgery system for lung cancer. *Expert review of medical devices*. 2012;9(5):465-75

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