TRANSITION TO THE RADIXACT® SYSTEM FOR PEDIATRIC TOTAL BODY IRRADIATION (TBI) TREATMENT

A 6-year-old boy presented with a high white blood cell count and a mediastinal mass. He was diagnosed with high-risk T-cell acute lymphoblastic leukemia (ALL) and treated in the ALLTogether1 trial¹. During induction therapy with regimen B, the patient continued to have high levels of minimal residual disease (MRD). Subsequently, the patient underwent consolidation treatment, leading to a decrease in MRD levels. As a result, the patient was recommended for a hematopoietic stem cell transplant (HSCT).

Case Challenge:

Due to the patient's high-risk disease, the HSCT was planned during his first complete remission. TBI was indicated as part of conditioning for stem cell transplant. This is the first patient treated since transitioning from the TomoTherapy® System to the Radixact® System. Pediatric TBI patients are treated on the Radixact System using TomoDirect™ to achieve a homogenous dose distribution and to minimize dose to the lungs.

FACTS AT-A-GLANCE



NHS Foundation Trust

FACILITY NAME

University Hospitals Birmingham NHS Trust

LOCATION

Birmingham, United Kingdom

ABOUT

The first pediatric TBI case treated using the Radixact® System, demonstrating workflow efficiencies

TREATMENT SYSTEM EXPERIENCE

Radixact X7 System

TECHNOLOGY SOLUTIONS

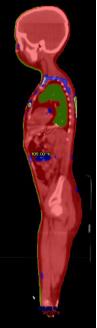
- VOLO® Ultra Optimizer
- PreciseART® Offline Adaptive Radiation Therapy Option
- ClearRT®

Treatment:

The patient was treated under general anesthesia with comprehensive immobilization using Medtech head curve Q2, Moldcare cushion, full body vac bag on lock board and foot stocks. Tattoos were placed anteriorly and laterally at the chest and pelvis for alignment. Helical kVCT images using ClearRT® were performed using a whole-body protocol with a field of view (FOV) of 500 mm.

The plan was created using VOLO® Ultra Optimizer in TomoDirect™ mode, with a fixed jaw and a 5 cm field. Ten fields were used to deliver a median dose of 12 Gy across six fractions, ensuring uniform dose distribution. The mean dose to the inner lungs was reduced to approximately 10.5 Gy. Special attention was given to





Planning CT scan

ClearRT® scan

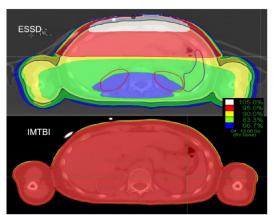
the spleen, included as an optimization target. Robustness plans were created to evaluate the effect of a 10 mm shift in all cardinal directions. Plan assessments were performed using PreciseART® Offline Adaptive Radiation Therapy option for the first two fractions.

Outcomes:

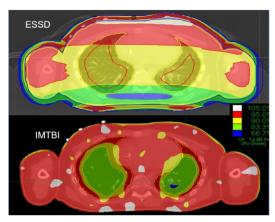
- No grade >2 side effects observed during treatment
- Patient discharged from hospital seven weeks after stem cell transplant



Rationale for Transitioning from Extended Source to Surface Distance (ESSD) Technique to the Radixact® System



		PTV (% OF PRESCRIBED)	
Volume %	Dose %	IMTBI	ESSD TBI
100 (99.96)	>90	92.0	78.2
50	100	100.2	90.5
0 (0.04)	>120	110.3	116.0
Large Volume Homogenity Index PTV	$\frac{D_{0.04} - D_{99.96}}{D_{50}}$	0.18	0.38



OAR	Dose Metric	Objective	Achieved	Achieved
			IMTBI	ESSD TBI
Spleen	Mean	12Gy	12.12	10.24
	D100	>95%	96.1	81.2

	OAR	Dose Metric	Objective	Achieved	Achieved
				IMTBI	ESSD TBI
	Inner Lungs	D _{50%}	10Gy	10.2	11.2
		V _{8Gy}	>95%	100	100

TomoTherapy® System Vs. Radixact System

	TomoTherapy® System	Radixact® System	Time saved (seconds)
Imaging tim (seconds)	896.00	101.40	794.60
Treatment tin (seconds)	e 1065.36	877.70	187.70

Conclusion:

- · Using the Radixact System, there was significant improvement in plan quality compared to the ESSD technique
- Implementing time reductions on Radixact System vs. TomoTherapy System has resulted in a more tolerable treatment experience and a decreased impact on the department workflow
- Assessments can be effectively performed using the PreciseART® Offline Adaptive Radiation Therapy software, ensuring the plan's robustness
- Treatment was tolerated well by the patient

Most side effects of radiotherapy, including radiotherapy delivered with Accuray systems, are mild and temporary, often involving fatigue, nausea, and skin irritation. Side effects can be severe, however, leading to pain, alterations in normal body functions (for example, urinary or salivary function), deterioration of quality of life, permanent injury, and even death. Side effects can occur during or shortly after radiation treatment or in the months and years following radiation. The nature and severity of side effects depend on many factors, including the size and location of the treated tumor, the treatment technique (for example, the radiation dose), and the patient's general medical condition, to name a few. For more details about the side effects of your radiation therapy, and to see if treatment with an Accurary product is right for you, ask your doctor. Accurary incorporated as a medical device manufacturer cannot and does not recommend specific treatment approaches. Individual results may vary.



¹ https://www.cancerresearchuk.org/about-cancer/find-a-clinical-trial/a-study-looking-at-treatment-children-young-adults-with-newly-diagnosed-acute-lymphoblatic-leukaemia-alltogether1 Important Safety Information: