

Multicenter implementation of a CT scanner dose excellence program based on clinical indication, BMI and diagnostic image quality assessment



H.Brat⁽¹⁾, S.Imsand⁽¹⁾, C.Dias⁽²⁾, C.Thouly⁽¹⁾, S.Montandon⁽³⁾, B.Rizk⁽¹⁾, D.Fournier⁽¹⁾, F.Zanca⁽²⁾

⁽¹⁾Groupe 3R, Switzerland, ⁽²⁾GE Healthcare, ⁽³⁾Philips Healthcare
No disclosure

Background

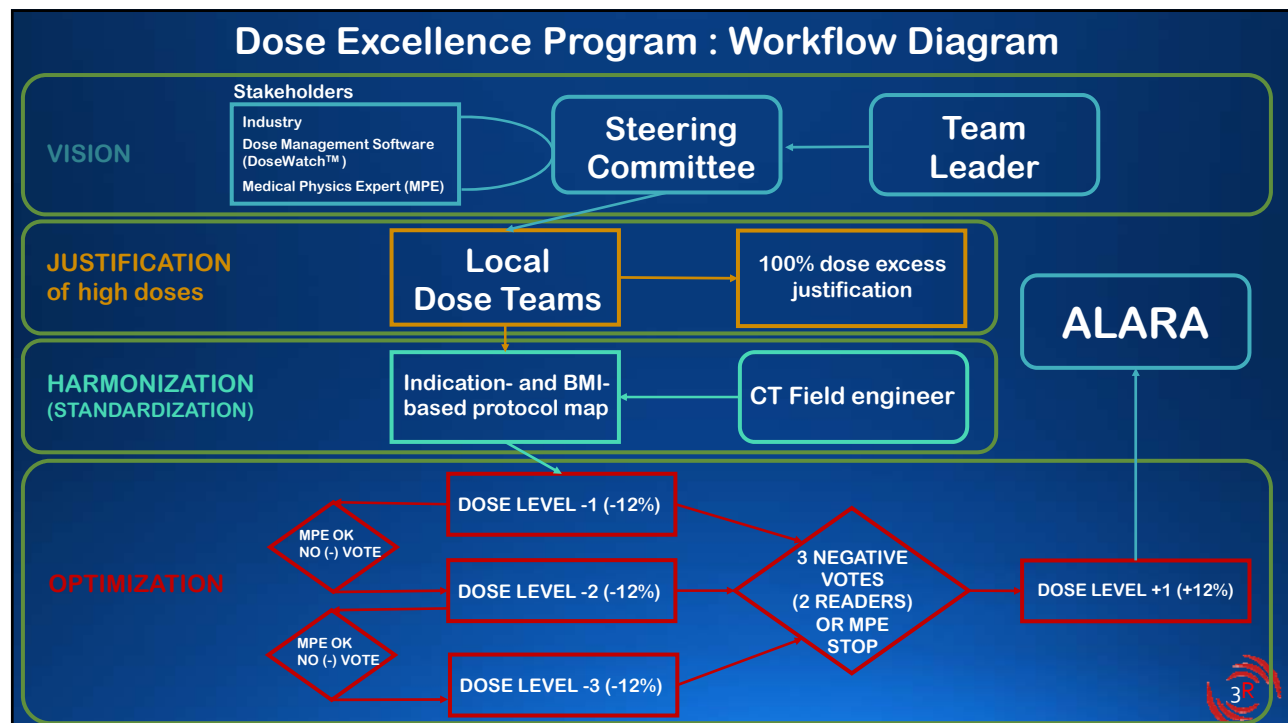
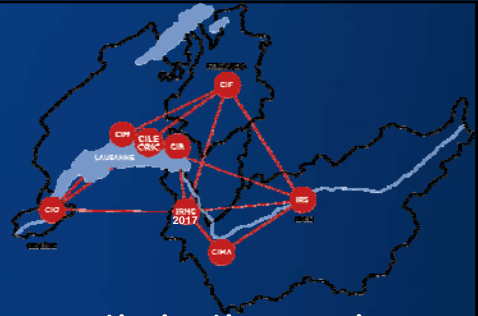
- The revised European Basic Safety Standards Directive (2018) requires :
 - the radiation dose of every CT exam to be recorded with investigation of cases where radiation dose exceeded established reference levels.
 - Radiation protection education and dose optimization training.
- Initial multicenter CT scanner data analysis in Groupe 3R identified :
 - large dose variations,
 - protocol parameters inhomogeneity,
 - lack of staff training uniformity.



Purpose

• Groupe 3R Board priorities (2016)

- To define and set-up a radiation dose optimization and education program at the group level (7 centers and 7 CT scanners from 3 manufacturers).
- To implement a “dose culture” by guiding 22 radiologists and 40 technologists towards a change of practice.



Step 1 : Harmonization phase

Radlex

Indication-based mapping



BMI < 25

BMI > 25

- Protocol Radlex mapping
- Design of a clinical indication-based protocol map
- 2 categories of patients for each protocol, according to BMI
- 30 protocols per BMI category



Step 1 : Harmonization phase

Acquisition parameters: Indication and BMI based

- Detector configuration
- Pitch
- Tube rotation time
- Tube voltage
- Tube current modulation
- Noise index
- Reconstruction kernel
- Reconstruction parameters
- Reconstruction standardization



Purpose

To deliver a $CTDI_{vol}$ value

- Close to P25 DRL when BMI < 25
- Below P75 DRL when BMI > 25



Diagnostic image quality assessment

Radiologists prospectively vote for diagnostic image quality using an electronic voting tool in the dose management software



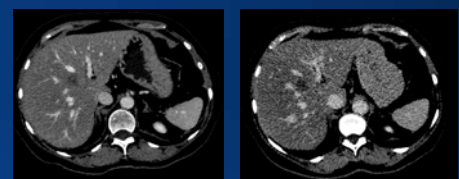
A negative vote needs to be confirmed by a second reader using adapted European image quality guidelines

	YES	No
Visually sharp reproduction of the liver parenchyma and intrahepatic portal veins	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the liver veins	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the structures of the liver hilus	<input type="radio"/>	<input checked="" type="radio"/>
Visually sharp reproduction of the common hepatic duct	<input type="radio"/>	<input checked="" type="radio"/>
Reproduction of the ductus choledochus (common bile duct) in the pancreatic parenchyma	<input type="radio"/>	<input checked="" type="radio"/>
Reproduction of the gallbladder wall	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the splenic artery	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the extrahepatic portal vein system including v. lumbalis and v. mesenterica sup.	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the origin of the coeliac trunk	<input checked="" type="radio"/>	<input type="radio"/>
Visually sharp reproduction of the mesenteric artery	<input checked="" type="radio"/>	<input type="radio"/>

Step 2 : Optimization phase

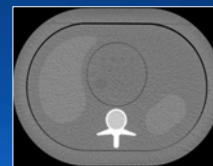
• Dose reduction phase

- 12% step-wise mA reduction for all protocols every 50 examinations of the same indication.
- In case of 3 negative voting for diagnostic image quality per protocol, confirmed by a second reader, dose was increased by 12% to reach previous accepted dose level (ALARA).
- In parallel, the Medical Physics Expert quantified with a model observer the low contrast detectability using an anthropomorphic phantom (QRM™ 401 abdomen phantom, Germany) to assess at which mA reduction level a 5mm lesion in the liver would not be diagnostically detectable.

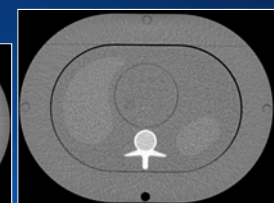


Before optimization

After optimization



Size M, correlated to BMI < 25*



Size L, correlated to BMI > 25*

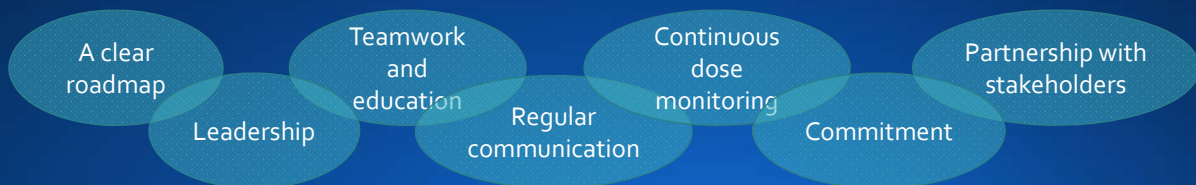
*Comparison of clinical and phantom image quality for low contrast liver lesions. SSJ22 Physics : CT Radiation Dose 1, Tuesday 3 PM, Room S403B



Results

- A think tank on image quality related to clinical indication and patient habitus enabled a team commitment in a quality project and a consensual good practice standardization.
- The use of a dose management software combined to Radlex protocol mapping enabled 100% of dose excess justification and protocol comparison in a multicenter setting.
- Protocol harmonization allowed comparison of comparable data (no redundancy), maintained diagnostic image quality and reduced dose by 6% for chest and 7% for abdomen.
- Protocol optimization enabled an additional average dose reduction of 26% (range 20-30%, depending on clinical indication), before hitting the low contrast resolution limit as assessed by phantom measurements.
- Clinical indication- and BMI-based protocols allowed significantly lower dose levels than existing DRL based on anatomical region with a sufficient diagnostic image quality : “The right dose for the right diagnosis”.

The complex process of
homogenizing CT protocols
and optimizing radiation doses
without compromising image quality
can be achieved with



hugues.brat@groupe3R.ch

