\*JSMP医学物理サマーセミナー2011;

閉講式 : 9/3:12:00~12:15 \*お知らせ

①セミナー講義資料は、以下よりログインし、必ず、事前に資料ダウンロ ードの上参加ください。LOG IN

②9/1;熊本空港⇒会場Hotel送迎バス;

·第1便;10:15発予定

·第2便:11:55発予定 送迎バス利用者集合場所・発射場所:

熊本空港; 団体ロビー⇒国際線ターミナルビル前 (MAP.pdf)

バスは国際線ビル前に停車しています。発車5分前には各自バスに乗車下 さい。\_\_\_バス利用希望など事前申込み者番号リスト

③9/1:ホテルで昼食を必要とする方

9月1日昼食はプログラムに含まれていません;ホテルでの昼食を希望す る場合は申込みが必要です。 申込み者リストを確認下さい。 基本的にホテ

ル昼食は予約制になっていますのでご注意ください。

ホテル昼食の追加申込みは事務局宛にご連絡下さい(〆切;8/23)

④他交通手段ご利用の場合は現地集合でお願い致します。 アクセス; http://www.aso-ikoi.jp/access/index.html 1) JR「阿蘇駅」、JR「いこいの村駅」到着の方はホテルへ電話連絡に

よりホテルからの送迎車が利用可能です。(tel:0967-34-2151) 2) 熊本空港からリムジンバスを利用される方は、阿蘇・大分方面行きを 利用し、JR「阿蘇駅」で下車してくだい。(tel:0967-34-2151)

⑤ 9/3: 会場Hotel⇒熊本空港行きバス ·第1 便:12:20 頃発—13:20 頃着予定 ・第2便:13:00過ぎ発

事務局:JSMP教育委員会サマーセミナー実行委員 mailto: <jsmp11 summer@nirs.go.jp> 問合せsubject名<Summer Seminar>

Medical Physics Summer Seminar 2011 in ASO-ikoinomura-kumamoto

## JSMP医学物理サマーセミナー2011

September 1-3,2011 Aso-KUMAMOTO

Day 1: Thursday Sep 1 Welcome, Course overwiew 13:15 13:00 Radiation Protection and Safety 1(赤羽恵一) 13:15 14:35 Radiation Protection and Safety 2(赤羽恵一) 14:45 16:05 Magnetic Resonance Imaging 1(山本 徹) 17:35 16:15 Magnetic Resonance Imaging 2(山本 徹) 17:45 19:05 Banquet 19:30

Walking 5:30 6:30 Breakfast 7:00 8:30 External Photon Beams: Physical Aspects(荒木不次男) 9:00 10:20 Electron Beams: Physical and Clinical Aspects(荒木不次男) 10:35 11:55 Lunch 12:00 13:00 Nuclear Medicine/Imaging 1(山谷泰賀) 14:00 15:20 Nuclear Medicine/Imaging 2(山谷泰賀) 16:55 15:35 18:00 19:00 Supper

Night session & Informal Q&A

Day 2: Friday Sep 2

Day 3: Saturday Sep 3 Breakfast 8:30 7:00 MV-CBCT(IGRT)(隅田伊織) 10:20 9:00 Special Procedures and Techniques in Radiotherapy(SRS,SRT)(塩見浩也) 10:35 11:55 Closing remark 12:00 \*本プログラム参加の機構認定単位数は下記の通り:

医学物理士認定機構; 医学物理士業績評価単位::10

放射線治療品質管理機構認定単位: C2-1

20:00

22:00

# Syllabus



#### 赤羽 恵一(放医研) 1. Introductions and Historical Perspective

1-1. Radiation Protection and Radiation Safety

(b) Observed radiation injury (c) Suggested radiation protection practices

(a) Discovery and early application of ionizing radiation

- (d) Pre-regulatory initiatives 2. Interaction Physics as Applied to Radiation Protection
- (a) Indirectly and directly ionizing radiation (b) Bethe-Bloch formalism for coulomb scattering, shell effects,
- polarization phenomena, nuclear processes, adiabatic scattering, track structure, target
- phenomena, radioactive processes, Anderson-Ziegler parameterization, Janni tabulation,
- and effects due to mixtures and compounds
- (c) Electromagnetic interaction: photoelectric effect, Compton effect, pair production,
- shower cascade phenomena
- (d) Neutron interactions: elastic and non-elastic processes
- 3. Operational Dosimetry (a) Units
- (b) Kerma and absorbed dose
- (c) Dose equivalent ii. Dose/dose equivalent interpretation
- ii. Pulse field response
- 5. Shielding: Properties and Design (a) Directly ionizing particles
- (b) Indirectly ionizing particles
- (c) Build-up parameterization (d) Stochastic sampling: Monte Carlo
- Source description and sampling
- ii. Interaction sampling iii. Geometry effects
- iv. Scoring
- v. Public domain codes (e) Particle Accelerators
- i. Primary particle shielding
- ii. Secondary-tertiary particle shielding iii. Energy and particle type dependence
- iv. Interlocks and access control v. Modeling radiation environment
- (f) NCRP (National Council on Radiation Protection and
- Measurements) shielding recommendations and techniques
- 6. Statistics (a) Statistical interpretation of instrument response
- (b) Design of experiments
- (c) Stochastic and nonstochastic error analysis (d) Interpreting experimental results
- 7. Radiation Monitoring of Personnel (a) Instrumentation and techniques
- (b) Integral and active devices (c) Dynamic range and response sensitivities
- (d) Film, TLD, Lexan, and CR-39
- (e) Pocket ion chambers and GM counters (f) Pregnant workers and fetal dose limits
- 8. Internal Exposure (a) ICRP 26, ICRP 2A recommendations
- (b) Medical internal radiation dose (MIRD) dosimetry (c) Monitoring and radiation control
- (d) Biological assay (e) Dispersion in a working environment (f) Allowed limit of intake and derived air (or water)
- concentrations 9. Environmental Dispersion
- (a) Release of radionuclides to the environment (b) Dosimetric consequences
- (c) Environmental Protection Agency (EPA) and U.S. Nuclear Regulatory Commission
- (NRC) air and water dispersion models 10. Biological Effects
- (a) Basic radiation biology (b) Nonstochastic and stochastic responses
- (c) Biological experimental data base of radiation injury
- (d) BEIR (Biological Effects of Ionizing Radiation) and UNSCEAR (United Nations Scientific Committee on the Effects of Atomic Radiation) Reports
- (e) Patient and fetal dose issues 11. Regulations (a) What is; what is not
- (b) 10CFR19-70; 49USDOT300-399, 198; 219SFDA 278;
- 290SHA; 42USPHS; 40USEPA
- (c) States: agreement or not (d) Relationship to NCRP and ICRP (International Commission on
- Radiation Protection) 12. High/Low Level Waste Disposal
- (a) USNRC/USDOE/USEPA Repository (U.S. Nuclear Regulatory Commission/
- Department of Energy/Environmental Protection Agency)
- (b) Low level compacts (c) Future impacts
- 13. Nonionizing Radiation (a) Electromagnetic and sound hazards
- (b) Device emission requirements (c) Measurement techniques

1-2 Magnetic Resonance Imaging

(d) Regulatory control

1. Basic Principles

### (a) Intrinsic and extrinsic parameters affecting MR image contrast (b) Required properties of nuclei that are useful in MR

(c) The static magnetic field (B<sub>0</sub>) and the equilibrium distribution (d) The Larmor frequency and the radiofrequency field (B1) (e) The lab and rotating frames of reference

山本 徹(北海道大学)

- (f) Relaxation mechanisms (T<sub>1</sub>, T<sub>2</sub>, T<sub>2</sub>\*) and effects of common contrast agents
- (g) The basic spin-echo sequence
- (h) Contrast in spin-echo imaging (i) Spatial encoding using linear magnetic field gradients ( $G_x$ ,  $G_y$ ,
- Gz) i. Slice selection
- ii. Frequency-encoding iii. Phase-encoding
- (j) Properties of "k-space" 2. Hardware
- (a) The static magnetic field subsystem i. Common field strengths and magnet designs

iv. 2D vs. 3D acquisitions

- ii. Siting issues (b) The radiofrequency (RF) field subsystem
- . Coil designs: volume, surface, phased array
- ii. Radiofrequency shielding requirements (siting) (c) The gradient field subsystem i. Maximum amplitudes, risetimes, and slew rates
- ii. Eddy current effects and compensation techniques 3. Basic Image Quality Issues
- (b) Resolution (c) Image acquisition time

- 4. Basic Pulse Sequences (a) Spin-echo sequence
- (b) Gradient-echo sequences
- (c) Fast spin-echo sequence (d) Inversion recovery sequences and applications [STIR, FLAIR
- (Short Time Inversion Recovery, Fluid-Attenuated Inversion Recovery)] (e) Common sequence options (spatial and chemical saturation
- techniques) (f) Ultrafast imaging sequences (echo planar imaging and spiral
- techniques) (g) MR flow sensitive sequences
- i. Flow-related phenoma ii. Time-of-flight MRA
- iii. Phase contrast MRA iv. Bolus contrast agent-enhanced MRA
- vi. Diffusion-weighted and diffusion tensor imaging

v. Perfusion sensitive imaging

ii. Imaging methods iii. Experiment design and analysis

(h) Functional MRI neuroimaging techniques

- (i) MR spectroscopy (MRS) sequences (j) Parallel imaging techniques
- (d) Chemical shift (e) Truncation

(f) System-related

iii. Ghosting

i. Physiological basis

- i. Distortions ii. RF coil problems and RF interference
- iv. Receiver/memory/array processor problems (g) Spatial accuracy limits and optimization

physiological effects)

control aspects

- 6. Safety and Bioeffects (a) Static field considerations (projectile, effects on implants,
- (c) Gradient field considerations (peripheral nerve stimulation, sound pressure levels)
- (d) Food and Drug Administration (FDA) guidelines (e) MR and pregnant patients, technologists, and nursing staff
- (a) The ACR (American College of Radiology) standards related to MRI (b) The ACR MR Accreditation Program (MRAP)
- (d) Other guidelines, including AAPM task group reports and NEMA (National Electrical Manufacturers Association) reports
- 2-1. External Beam Radiation Therapy 荒木不次男 (熊本大学) 1. Clinical Photon Beams: Description
- (a) Basic parameters: Field size, source-skin distance, source-axis distance, sourcecollimator distance
- (c) Field collimators: Primary, secondary, and tertiary placement of collimators;
- rectangular (upper and lower jaws); circular; multileaf collimators 2. Clinical Photon Beams: Point Dose Calculations (a) Percentage depth dose (PDD)
- (b) Peak-scatter factor (PSF) (c) Tissue-air ratio (TAR)
- (f) Scatter function (g) Scatter-air ratio (SAR)
- (h) Scatter-maximum ratio (SMR) (i) Collimator factor (j) Relative dose factor/output factor
- (a) Factors affecting the fundamental dosimetry quantities (b) Relationships between the fundamental dosimetry quantities (c) Collimator and phantom scatter corrections
- (d) Irregular fields and Clarkson's integration method
- (a) Electron treatment head i. Energy selection
- ii. Beam broadening methods: dual scattering foil vs. scanned beam iii. Collimating methods: trimmers vs. applicators (cones)
- (c) Energy spectrum i. Characteristics (\_E, E<sub>p</sub>)
- (d) Dose distribution i. Beam flatness and symmetry
- ii. Penumbra iii. Isodose plots
- ii. Output factor formalisms (f) Effect of air gap on beam dosimetry

(g) Fundamental principles

- i. Square-root method ii. Effective vs. virtual source
- 5. Special Photon and Electron Beams
- i. Linacs with multileaf collimators
- ii. Tomotherapy iii. Stereotactic beams and robotic linacs (b) Intensity-modulated radiation therapy with electron beams

2-2. Nuclear Medicine/Imaging 山谷泰賀 (放医研)

# 1. The Gamma Camera

- (a) Camera characteristics (b) Collimators
- (c) Crystals (d) Photomultiplier tube array (e) Image formation
- (g) The pulse height analyzer 2. Radionuclide Image Quality
- (b) Blur and visibility of detail (c) Image noise

(a) Contrast

modes

(f) Spectrometry

- ii. Clinical PET imaging procedures
- . Principles of SPECT imaging, hardware, resolution ii. Clinical SPECT imaging procedures
- 5. Patient Exposure and Protection (a) Internal dosimetry
- procedures
- 6. Personnel Exposure and Protection
- (c) Exposure sources
- (f) Exposure from radioactive sources
- (b) Survey meters (c) Activity measurement
- 8. Principles of Radiochemistry, Radioimmunoimaging, and the Radiopharmacy
- (b) Radioimmunoimaging and radioimmunotherapy principles (c) Radiopharmacy techniques 9. Quality Control Issues in Nuclear Medicine

#### 隅田伊織 (大阪大学) 3.1.1 Portal Imaging

3-1.IMAGING FOR TREATMENT GUIDANCE AND

(e) Registration to DRR (f) Imaging dose (g) QA

(d) DRR calculation from CT

MONITORING

- 3.1.2 Cone-Beam CT (a) Large-field CT, field size
- (e) Imaging artifacts (g) QA
- 1. Special External Beam Radiotherapy Techniques: Basic Characteristics, Historical Development, Quality Assurance (Equipment and Treatment),

塩見浩也 (大阪大学)

3.2 Special Techniques in Radiotherapy

- (a) Stereotactic radiosurgery (b) Stereotactic radiotherapy

- (d) Uniformity (e) Clinical gamma camera applications 3. Radionuclide Tomographic Imaging (a) Positron Emission Tomography (PET) and PET-CT . Principles of PET imaging, hardware, resolution, acquisition iii. Quantitative PET imaging iv. Cine (4D) PET (b) Single Photon Emission Computed Tomography (SPECT)
- iii. Quantitative SPECT imaging 4. Statistics: Counting Error
- (b) Clinical dosimetry and typical doses for common imaging (c) Radionuclide therapy dosimetry
- (a) Effective dose equivalents (b) Exposure limits
- (d) Area shielding (e) Personnel shielding
- 7. Radiation Measurement (a) Ionization chambers
- (a) Radiochemistry principles
- (c) The ACR MR Quality Control Manual and its recommended quality

#### (a) Portal film, electronic portal imaging (b) Types of imaging panels, technologies (c) Scatter

- (b) MV cone-beam CT (c) Scatter, scatter rejection (d) Imaging quality (f) Imaging dose (Dose incorporation to plan)
- Diseases Treated

- (a) Intensity-modulated radiation therapy with photon beams

- 5. Artifacts and Methods for Artifact Rejection/Reduction iii. TLD energy, dose, dose rate response (a) Motion (c) Dose equivalent instrumentation (b) Aliasing or "wrap-around" . Energy dependence (c) Metal objects
  - (b) RF field considerations (tissue heating, specific absorption rate, burn injuries)
  - (f) Common MR contrast agents 7. Quality Control

  - (b) Field size options: Circular, square, rectangular, irregular
  - (d) Tissue-maximum ratio (TMR) (e) Tissue-phantom ratio (TPR)
  - (k) Off-axis ratio 3. Clinical Photon Beams: Basic Clinical Dosimetry
  - (e) Tissue heterogeneities and corrections 4. Clinical Electron Beams
  - (b) Depth-dose distribution i. Characteristics (Ds,Dx,R100,R90,Rp,R90-10) ii. Variation with energy and field size
  - ii. Specification at surface (range-energy relationships) and depth
  - i. Method of dose prescription

(e) Determination of monitor units

- iii. Side-scatter equilibrium

- (a) Signal-to-noise ratio and contrast-to-noise ratio in MRI

- 会期:9月1日(木)~9月3日(土) 会場受付:9/1:12:30-13:00 開講式 : 9/1:13:00~13:15
- 会場: <u>リゾートホテル阿蘇いこいの村</u> (tel:0967-34-2151)