DISCOVERY MI PET/CT
FINAL STUDY

GE Healthcare

TYPICAL

A mandatory component of this drawing set is the GE Healthcare Pre Installation manual. Failure to reference the Pre Installation manual will result in incomplete documentation required for site design and preparation. Pre Installation documents for GE Healthcare products can be accessed on the web at: www.gehealthcare.com/siteplanning

GE does not take responsibility for any damages resulting from changes on drawings made by others. Errors may occur by not referring to the complete set of final issue drawings. GE cannot accept responsibility for any damage due to the partial use of GE final issue drawings, however caused. All dimensions are in millimeters unless otherwise specified. Do not scale from printed pdf files. GE accepts no responsibility or liability for defective work due to scaling from these drawings.
GENERAL SPECIFICATIONS

- GE is not responsible for the installation of developers and associated equipment, lighting, cassette trays and protective screens or derivatives not mentioned in the order.
- The final study contains recommendations for the location of GE equipment and associated devices, electrical wiring and room arrangements. When preparing the study, every effort has been made to consider every aspect of the actual equipment expected to be installed.
- The layout of the equipment offered by GE, the dimensions given for the premises, the details provided for the pre-installation work and electrical power supply are given according to the information noted during on-site study and the wishes expressed by the customer.
- The room dimensions used to create the equipment layout may originate from a previous layout and may not be accurate as they may not have been verified on site. GE cannot take any responsibility for errors due to lack of information.
- Dimensions apply to finished surfaces of the room.
- Actual configuration may differ from options presented in some typical views or tables.
- If this set of final drawings has been approved by the customer, any subsequent modification of the site must be subject to further investigation by GE about the feasibility of installing the equipment. Any reservations must be noted.
- The equipment layout indicates the placement and interconnection of the indicated equipment components. There may be local requirements that could impact the placement of these components. It remains the customer’s responsibility to ensure that the site and final equipment placement complies with all applicable local requirements.
- All work required to install GE equipment must be carried out in compliance with the building regulations and the safety standards of legal force in the country concerned.
- These drawings are not to be used for actual construction purposes. The company cannot take responsibility for any damage resulting therefrom.

CUSTOMER RESPONSIBILITIES

- It is the responsibility of the customer to prepare the site in accordance with the specifications stated in the final study. A detailed site readiness checklist is provided by GE. It is the responsibility of the customer to ensure all requirements are fulfilled and that the site conforms to all specifications defined in the checklist and final study. The GE Project Manager of Installation (PMI) will work in cooperation with the customer to follow up and ensure that actions in the checklist are complete, and if necessary, will aid in the rescheduling of the delivery and installation date.
- Prior to installation, a structural engineer of record must ensure that the floor and ceiling is designed in such a way that the loads of the installed system can be securely borne and transferred. The layout of additional structural elements, dimensioning and the selection of appropriate installation methods are the sole responsibility of the structural engineer. Execution of load bearing structures supporting equipment on the floor, wall or walls is the customer’s responsibility.

RADIO-PROTECTION

- Suitable radiological protection must be determined by a qualified radiological physicist in conformance with local regulations. GE does not take responsibility for the specification or provision of radio-protection.

THE UNDERSIGNED, HEREBY CERTIFIES THAT I HAVE READ AND APPROVED THE PLANS IN THIS DOCUMENT.

<table>
<thead>
<tr>
<th>DATE</th>
<th>NAME</th>
<th>SIGNATURE</th>
</tr>
</thead>
</table>

DISCLAIMER

RADIO-PROTECTION

CUSTOMER RESPONSIBILITIES

GLOBAL SITE READINESS CHECKLIST (DI)

<table>
<thead>
<tr>
<th>Customer Name:</th>
<th>PMI Name:</th>
</tr>
</thead>
<tbody>
<tr>
<td>GON/SO Number:</td>
<td>Field Service Name:</td>
</tr>
<tr>
<td>Equipment:</td>
<td>Country/City or City/State:</td>
</tr>
<tr>
<td>Site Visit Date for SRC:</td>
<td>SRC Status:</td>
</tr>
</tbody>
</table>

Site Ready Checks at Installation

General Site Planning

- Room dimensions, including ceiling height, for all Exam, Equipment/Technical & Control rooms meets GE specifications.
- Ceiling support structure, if on the GE drawing, is at correct location and height according to the drawing specifications. Levelness and spacing has been measured. Overhead support Structure has been confirmed with contractor to meet GE criteria.
- Rooms that will contain equipment, including staging areas if applicable, are construction debris free. Precautions must be taken to prevent debris from entering rooms containing equipment.
- Delivery route from truck to installation space has been reviewed, all communications have occurred, arrangements made for special handling (if needed). Floors along delivery route will support weight of the equipment, reinforcements arranged if needed.
- System power & grounding (PDB/MDP) is available as per GE specifications, installed at point of final connection and ready to use. Lock Out Tag Out is available.
- System power and ground audit has been scheduled to be completed during installation of equipment. (If Required) GEHC PM to confirm if needed.
- Adequate room illumination installed and working.
- Cableways (floor, wall, ceiling, etc.) ready for GE cables and are of correct length and diameter. Cableways routed per GE Final drawings and access openings installed as determined by GEHC PM. Surface floor duct installed at time of system installation.
- HVAC systems Installed, and the site meets minimum environmental operational system requirements.
- Network outlets installed and computer network available and working.
- Hospital/IT connectivity contacts have been engaged and information has been added to Project management tool. (If Required)
- Floor levelness/flatness is measured and within tolerance, and there are no visible defects per GEHC specifications. Floor Strength and thickness have been discussed with customer/contactor and they have confirmed GE requirements are met.
- Customer supplied countertops where GE equipment will be installed are in place.

Specific for PET and Nuclear Medicine

- Nuclear Medicine systems levelness measurement survey must be provided to GE prior the delivery.
- Site has license for using/Importing radioactive sources and a Hot Lab is available. Radioactive Sources should be available for system calibration during installation.
- Doors and windows complete or scheduled to be installed. If applicable, radiation protection (shielding) finished & radioprotection regulatory approval for installation obtained.

PMI Signature:

Customer Signature:

FS Signature: optional

DOC1809666 Rev. 6
**ENVIRONMENT**

**ALTITUDE**
- Operating altitude: from -150 m [-492 ft] (below) to 2400 m [7875 ft] (above) sea level.

**MAGNETIC FIELD SPECIFICATIONS**
- Limit the magnetic interference to guarantee specified imaging performance.

**GANTRY**
- Ambient static magnetic fields less than 1 Gauss.
- Ambient AC magnetic fields less than 0.01 Gauss.

**OPERATOR CONSOLE**
- Ambient static magnetic fields less than 10 Gauss.

**MAXIMUM GANTRY AUDIBLE NOISE LEVEL**
- The maximum ambient noise level is produced by the gantry during a CT scan acquisition.
- It is less than 70 dBA when measured at a distance of one meter from the nearest gantry surface, in any direction.

**BACKGROUND RADIATION**
- It is important that background radiation be kept to a minimum. The coincidence detection used in a PET system allows a moderate amount of external singles events. The PET/CT system has been found to have less than 1% deadtime if the external field is below 1 mR/hr from a single source.
- Because area background can be more general than a single source, a lower limit is appropriate. If the area dose rate is maintained to less than 0.2 mR/hr (due to 511 or lower energy gamma rays) at the covers, detector deadtime should not exceed 1%.
- Radioactive sources must be stored in approved shielded containers. It is recommended that any radioactive source not specifically designed to be housed in the gantry's lead storage container be stored in a separate room (hot lab) adjacent to, and accessible from, the Scan Room. Doses should be prepared in the same area.

**VIBRATION SPECIFICATIONS**
- Shock Restrictions: The system cannot tolerate shock or vibration. System components cannot be tipped, dropped, or hoisted.
- The scanning facility shall be isolated from vibration such as: hospital power plants, pumps, motors, air handling equipment, air conditioning units, nearby rooms with exercise equipment or where exercise occurs, hallway foot traffic, elevators, parking lots, roads, subways, trains, and heliports; otherwise, vibration will affect the image quality of the scanner.
- CT systems are sensitive to vibration and may display limited performance if exceeding the vibration limits listed below. The band of frequencies in which systems exhibit the most sensitivity appears at or near the resonant frequencies of the gantry and the patient table, the latter of which varies depending on patient mass and location. These frequencies fall within the following ranges:
  - Patient Table: 2 – 10 Hz
  - Gantry: 8 – 14 Hz
- It is the customer’s responsibility to contract a vibration consultant or qualified engineer to verify that these specifications are met and implement an appropriate solution.
- The maximum steady state vibration transmitted through the floor should not exceed 2.5 mm/s² RMS maximum single frequency above ambient baseline from 0.5 to 80 Hz (measured in any 1 hour during a normal operating period).

**IMPORTANT CUSTOMER READINESS ALERT**

- This equipment involves the use of radioactive isotopes, including those sources necessary for equipment calibration. Appropriate regulatory compliance and licensing must be arranged by the customer early in the planning process and then demonstrated/available for equipment installation.
- Note: delivery path down corridors for gantry's and table must be evaluated prior to construction, as 90 degree turns require specific corridor width.

**CUSTOMER SITE READINESS REQUIREMENTS**

- Any deviation from these drawings must be communicated in writing to and reviewed by your local GE Healthcare Installation project manager prior to making changes.
- Make arrangements for any rigging, special handling, or facility modifications that must be made to deliver the equipment to the installation site. If desired, your local GE Healthcare Installation project manager can supply a reference list of rigging contractors.
- New construction requires the following:
  1. Secure area for equipment,
  2. Power for drills and other test equipment,
  3. Capability for image analysis,
  4. Restrooms.
- Provide for refuse removal and disposal (e.g. crates, cartons, packing)
- It is the customer’s responsibility to contract a vibration consultant/engineer to implement site design modifications to meet the GE vibration specification. Refer to the system preinstallation manual for the vibration specification.
## EXAM ROOM

### Technical Room Dimensions
- Finished Ceiling Height: 9'-0"
- Minimum door opening for equipment delivery: 55 in. w x 80 in. h [1397mm x 2032mm], contingent on a 96 in. [2438mm] corridor width

### Equipment Layout

#### BY ITEM

<table>
<thead>
<tr>
<th>ITEM</th>
<th>DESCRIPTION</th>
<th>MAX HEAT OUTPUT (btu)</th>
<th>WEIGHT (lbs)</th>
<th>MAX HEAT OUTPUT (W)</th>
<th>WEIGHT (kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A 1</td>
<td>Discovery MI Gantry</td>
<td>28320</td>
<td>6383</td>
<td>8300</td>
<td>2895</td>
</tr>
<tr>
<td>A 2</td>
<td>Patient Table</td>
<td>1023</td>
<td>1808</td>
<td>300</td>
<td>822</td>
</tr>
<tr>
<td>A 3</td>
<td>Power Distribution Unit</td>
<td>3400</td>
<td>813</td>
<td>1000</td>
<td>370</td>
</tr>
<tr>
<td>A 4</td>
<td>PARC4</td>
<td>6824</td>
<td>540</td>
<td>2000</td>
<td>246</td>
</tr>
<tr>
<td>A 5</td>
<td>Annulus Phantom Safe</td>
<td>-</td>
<td>-</td>
<td>149</td>
<td></td>
</tr>
<tr>
<td>A 6</td>
<td>Service Cabinet</td>
<td>-</td>
<td>90</td>
<td>-</td>
<td>41</td>
</tr>
<tr>
<td>A 7</td>
<td>Chiller</td>
<td>13649</td>
<td>320</td>
<td>4000</td>
<td>145</td>
</tr>
<tr>
<td>A 8</td>
<td>Power Distribution Box</td>
<td>-</td>
<td>66</td>
<td>-</td>
<td>30</td>
</tr>
<tr>
<td>A 9</td>
<td>Operators Console</td>
<td>3625</td>
<td>144</td>
<td>1100</td>
<td>65</td>
</tr>
<tr>
<td>D/B 10</td>
<td>Main Disconnect Panel</td>
<td>-</td>
<td>46</td>
<td>-</td>
<td>21</td>
</tr>
<tr>
<td>D 11</td>
<td>Injector - ceiling mounted</td>
<td>-</td>
<td>79</td>
<td>-</td>
<td>36</td>
</tr>
<tr>
<td>D 12</td>
<td>Injector Control and Electronics</td>
<td>320</td>
<td>22</td>
<td>94</td>
<td>10</td>
</tr>
<tr>
<td>D 13</td>
<td>Operators Chair</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>C 14</td>
<td>Counter top for equipment</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>C 15</td>
<td>Counter top with sink, base and wall cabinets</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>C 16</td>
<td>Lead glass window</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>C 17</td>
<td>Minimum door opening for equipment delivery: 55 in. w x 80 in. h [1397mm x 2032mm], contingent on a 96 in. [2438mm] corridor width</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Technical Room

- 11'-10" x 27'-10"

### Exam Room Height

- Finished Ceiling Height: 9'-0"
RADIATION PROTECTION LAYOUT

SHIELDING REQUIREMENTS:
Engage a qualified radiological health physicist to review your scan room shielding requirements, taking into consideration:

- Scatter radiation levels within the scanning room.
- Equipment placement.
- Weekly projected work-loads (number of patients/day technique (kVp*mA)).
- Materials used for construction of walls, floors, ceiling, doors, and windows.
- Activities in surrounding scan room areas.
- Equipment in surrounding scan room areas (e.g., film developer, film storage).

For small and medium filter survey, the 20 cm water phantom should be placed on the phantom headholder inserted into the end of the patient table.

The four scatter surveys depict measured radiation levels within the scanning room at the indicated distances, while scanning a 16 cm CTDI phantom for the Head Scan mode and 32 cm CTDI phantom for the Body Scan Mode. Use the mA, kV and aperture scaling factors in the table shown here to adjust exposure levels to the scan technique used at the site.

For example: The exposure level for a 120 kV, 800 mA, 1 sec scan at 50” (127 cm) away from the scan plane is:

\[
\text{exposure level} = 10.4 \mu\text{Gy} \times 0.71 \times \frac{800}{100} = 59.1 \mu\text{Gy}
\]

NOTE: Actual measurements can vary. Expected deviations equal ±15%, expect for the 5 mA and 1.25mm techniques, where variations may be greater (up to a factor of 2), due to the inherent deviation in small values. The maximum deviation anticipated for tube output equals ±40%.

RADIOACTIVE ISOTOPES

RADIOACTIVE ISOTOPES AND RADIOPROTECTION

Since the system produces X-ray radiation and involves the use of radioactive isotopes, compliance with Nuclear Regulatory Commission regulations (or country similar regulatory requirements), must be adhered to and all permissions obtained well in advance.

It is Customer's responsibility consult a qualified radiological health physicist for radiation protection requirements for the walls, floor, ceiling, doors, window glass, etc. (lead content and thickness) and warning lights and signs, in accordance with local requirements.

It is essential that regulatory compliance and preparations are completed early so that required source materials can be obtained prior to installation, including calibration sources and isotopes. These sources and isotopes may have fairly long delivery lead times and a short half-life, so that it may not be advisable to store them over long periods of time.

RADIOACTIVE SOURCE - ISOTOPE

The PET/CT system uses one radioactive source during calibration and the Daily QA Check.

<table>
<thead>
<tr>
<th>Isotope</th>
<th>Activity level</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ge-68</td>
<td>55 MBq ± 20%</td>
</tr>
<tr>
<td>Fluorine 18</td>
<td></td>
</tr>
<tr>
<td>Carbon 11</td>
<td></td>
</tr>
<tr>
<td>Nitrogen 13</td>
<td></td>
</tr>
<tr>
<td>Oxygen 15</td>
<td></td>
</tr>
</tbody>
</table>

Typical Positron Emitting Isotopes include

It is customer's responsibility provide isotopes for system calibration and prepare the required doses.
**SHIPPING DOLLY DIMENSIONS FOR GANTRY**

**CT GANTRY**

- **Weight with dollies and side rails = 2050 kg** [4520 lbs]

**PET IMAGE RING FOR WELDMENT GANTRY**

- **Weight with dollies and side rails = 1204 kg** [2654.3 lbs]

**DELIVERY**

- **THE CUSTOMER/CONTRACTOR SHOULD:**
  - Provide an area adjacent to the installation site for delivery and unloading of the GE equipment.
  - Ensure that the dimensions of all doors, corridors, ceiling heights are sufficient to accommodate the movement of GE equipment from the delivery area into the definitive installation room.
  - Ensure that access routes for equipment will accommodate the weights of the equipment and any transportation, lifting and rigging equipment.
  - Ensure that all necessary arrangements for stopping and unloading on public or private property not belonging to the customer have been made.

**DIMENSIONS OF DELIVERY WITH DOLLY TRANSPORT EQUIPMENT**

<table>
<thead>
<tr>
<th></th>
<th>mm</th>
<th>in</th>
<th>kg</th>
<th>lbs</th>
</tr>
</thead>
<tbody>
<tr>
<td>CT GANTRY</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LENGTH</td>
<td>2810</td>
<td>111</td>
<td>2050</td>
<td>4520</td>
</tr>
<tr>
<td>WIDTH</td>
<td>1290</td>
<td>51</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HEIGHT</td>
<td>2000</td>
<td>79</td>
<td></td>
<td></td>
</tr>
<tr>
<td>PET WELDMENT GANTRY</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LENGTH</td>
<td>2794</td>
<td>110</td>
<td>1204</td>
<td>2654.3</td>
</tr>
<tr>
<td>WIDTH</td>
<td>1118</td>
<td>44</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HEIGHT</td>
<td>1880</td>
<td>74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>PATIENT TABLE</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LENGTH</td>
<td>3836</td>
<td>151</td>
<td>1241</td>
<td>2736</td>
</tr>
<tr>
<td>WIDTH</td>
<td>864</td>
<td>34</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HEIGHT</td>
<td>1410</td>
<td>55.5</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Above dimensions shown with side rails on. The minimum unobstructed hallway width is 1803 mm, the minimum clear doorway openings is 1067 mm to accommodate delivery of the system.
• Methods of support for the steelwork that will permit attachment to structural steel or through bolts in concrete construction should be favored. Do not use concrete or masonry anchors in direct tension.

• All units that are wall mounted or wall supported are to be provided with supports where necessary. Wall supports are to be supplied and installed by the customer or his contractors. See plan and detail sheets for suggested locations and mounting hole locations.

• All ceiling mounted fixtures, air vents, sprinklers, etc. To be flush mounted, or shall not extend more than 6,35mm (1/4") below the finished ceiling.

• Floor slabs on which equipment is to be installed must be level to 6.00mm (1/4") in 3050mm (10'-0")

• Dimensions are to finished surfaces of room.

• Customers contractor must provide all penetrations in post tension floors.

• Customers contractor must provide and install any non-standard anchoring. Documents for standard anchoring methods are included with GE equipment drawings for geographic areas that require such documentation.

• Customers contractor must provide and install hardware for "through the floor" anchoring and/or any bracing under access floors. This contractor must also provide floor drilling that cannot be completed because of an obstruction encountered while drilling by the GE installer such as rebar etc.

• It is the customer’s responsibility to perform any floor or wall penetrations that may be required. The customer is also responsible for ensuring that no subsurface utilities (e.g., electrical or any other form of wiring, conduits, piping, duct work or structural supports (i.e. post tension cables or rebar)) will interfere or come in contact with subsurface penetration operations (e.g. drilling and installation of anchors/screws) performed during the installation process. To ensure worker safety, GE installers will perform surface penetration operations only after the customer’s validation and completion of the “GE surface penetration permit”
<table>
<thead>
<tr>
<th>ITEM</th>
<th>DESCRIPTION</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Floor contact area for discovery gantry and patient table. See detail on Structural Detail sheets for more information.</td>
</tr>
<tr>
<td>2</td>
<td>Structural supports for fastening the overhead counterpoised suspension. Support should run continuous with no fittings extending below face of channel, be parallel, square, and in the same horizontal plane, above finished ceiling. Ensure mounting surface is installed level or plumb within +/- 1 degree, and is structurally sufficient to maintain a level or plumb condition under 110 lb (50kg) system load and maximum system moment of 4400 in-lb (500n-m). Methods of support that will permit attachment to structural steel or through bolts in concrete construction should be favored. Do not use screw anchors in direct tension. 14&quot; x 14&quot; x 1/2&quot; thick steel plate provided by manufacturer. See detail on structural detail sheets.</td>
</tr>
<tr>
<td>3</td>
<td>Support Backing, locate as shown</td>
</tr>
</tbody>
</table>
ANCHORING/LOADING DISTRIBUTION TO THE FLOOR

GE SUPPLIED GANTRY ANCHORS (2106573)

ANCHORING AND FLOOR REQUIREMENTS

FINISHED FLOOR REQUIREMENTS

- Installation requires a finish floor in the scan and control rooms.
- The floor surface in the scan room directly under the gantry and table must be level.
- The floor shall be no greater than 6 mm [0.25 in] out of level over a 3048 mm [120 in] range, with level defined as the horizontal surface between the highest and lowest points.
- The floor shall have a minimum concrete thickness of 127 mm [5 in].
- Shims should not be used to compensate for a floor that does not meet this requirement.
- These requirements apply to all installation types.

NOTES:

- If the concrete floor has a floor covering installed over it (such as floor tile), 17 or more openings 101.6 mm [4 in] in diameter will be cut into the floor covering to ensure the table and gantry rest on the concrete. (Openings are cut during installation.)

SCALE 1:25

1. PET primary scan plane axis
2. CT scan plane axis
3. Longitudinal axis
4. Cable and hose access
5. Cable access only
6. 5 anchoring points for the Table
7. 8 anchoring points for the Gantry
8. Alternative anchoring points

Center of gravity
Minimum anchor embedment: 89 mm (2106573)
**MEDRAD MOUNTING DETAILS FOR CEILING INJECTOR**

### Structural Supports
- (Customer/contractor supplied)
- 356mm x 356mm x 12.7mm [14 in x 14 in x 1/2 in] Steel plate provided by GE and installed by customer/contractor
- 9.5 mm [0.375 in] Bolts (not by GE)

### Mounting Assembly
- 3200mm-3353mm [126 in-132 in] to finished floor requires long post
- 3048mm [120 in] to finished floor requires intermediate post
- 2743mm-2896mm [108 in-114 in] to finished floor requires short post

### Scale 1:10

### Recommended height to floor
2083mm - 2235mm [82 in - 88 in]

### Not to Scale

**Verify mounting assembly dimensions with injector manufacturer.**

**Post** | **Lengths** | **GE ECat No.**
---|---|---
Short | 22.8 | S80 EBD07NB, ND, NG, NY, PB & PE
Intermediate | 33.5 | 810 EBD07PP, PN, PK, PR & PY
Long | 39.4 | 1000 EBD07NE, NF, NI, NN, PD & EBD18AA

**Max. Arc**
- 1660 [65.4 in]

**Min. Plate Height** | **Max. Plate Height**
---|---
Short | 108 | 2743 | 114 | 2896
Intermediate | 120 | 3048 | 132 | 3333
Long | 126 | 3200 | 132 | 3333

**Top of mounting assembly**

**Highest head position**

**Lowest head position**

**Injector**
### TEMPERATURE AND HUMIDITY SPECIFICATIONS

#### IN-USE CONDITIONS

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exam Room</td>
<td>PET Gantry</td>
<td>2.8</td>
<td>9554</td>
</tr>
<tr>
<td></td>
<td>CT Gantry</td>
<td>5.5</td>
<td>18766</td>
</tr>
<tr>
<td></td>
<td>Patient table</td>
<td>0.3</td>
<td>1024</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>9</td>
<td>29344</td>
</tr>
</tbody>
</table>

Exam room or Technical room*

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exam room</td>
<td>Power distribution unit (CT PDU)</td>
<td>1.0</td>
<td>3400</td>
</tr>
<tr>
<td></td>
<td>PARC 4 (Reconstruction Cabinet)</td>
<td>2.0</td>
<td>6824</td>
</tr>
<tr>
<td></td>
<td>Chiller</td>
<td>4.0</td>
<td>13649</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>7</td>
<td>23873</td>
</tr>
</tbody>
</table>

Exam room or Technical room*

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control Room</td>
<td>Operator console</td>
<td>0.84</td>
<td>2860</td>
</tr>
<tr>
<td></td>
<td>LCD Monitor (2 units, 170 BTU/50 Watts each)</td>
<td>0.1</td>
<td>340</td>
</tr>
<tr>
<td></td>
<td>Peripheral Media Tower (PMT)</td>
<td>0.13</td>
<td>425</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>1</td>
<td>3625</td>
</tr>
</tbody>
</table>

*Technical Room is not mandatory, the placements of these elements are recommended in the Exam Room.

**WARNING**

This chart contains only the principal components of the PET/CT system and does not include information about non-GE supplied equipment.

### STORAGE CONDITIONS

<table>
<thead>
<tr>
<th>Condition</th>
<th>Specification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Temperature</td>
<td>0°C to +30°C +32°F to +86°F</td>
</tr>
<tr>
<td>Relative humidity (1)</td>
<td>≤ 70% RH</td>
</tr>
<tr>
<td>Temperature gradient</td>
<td>≤ 3°C/h</td>
</tr>
<tr>
<td>Humidity gradient</td>
<td>≤ 5% RH/h</td>
</tr>
</tbody>
</table>

Material should not be stored for more than 6 month.

(1) Non-condensing

(2) Actual heat output is site specific and dependent on the specific configuration and customer usage.

### AIR RENEWAL

According to local standards. The HVAC system should be designed to provide 5 air changes per hour to maintain adequate air quality and temperature.

**NOTE**: In case of using air conditioning systems that have a risk of water leakage it is recommended not to install it above electric equipment or to take measures to protect the equipment from dropping water.

### HEAT DISSIPATION

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exam Room</td>
<td>PET Gantry</td>
<td>2.8</td>
<td>9554</td>
</tr>
<tr>
<td></td>
<td>CT Gantry</td>
<td>5.5</td>
<td>18766</td>
</tr>
<tr>
<td></td>
<td>Patient table</td>
<td>0.3</td>
<td>1024</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>9</td>
<td>29344</td>
</tr>
</tbody>
</table>

Exam room or Technical room*

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exam room</td>
<td>Power distribution unit (CT PDU)</td>
<td>1.0</td>
<td>3400</td>
</tr>
<tr>
<td></td>
<td>PARC 4 (Reconstruction Cabinet)</td>
<td>2.0</td>
<td>6824</td>
</tr>
<tr>
<td></td>
<td>Chiller</td>
<td>4.0</td>
<td>13649</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>7</td>
<td>23873</td>
</tr>
</tbody>
</table>

Exam room or Technical room*

<table>
<thead>
<tr>
<th>ROOM</th>
<th>DESCRIPTION</th>
<th>Max (kW)</th>
<th>Max (btu)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control Room</td>
<td>Operator console</td>
<td>0.84</td>
<td>2860</td>
</tr>
<tr>
<td></td>
<td>LCD Monitor (2 units, 170 BTU/50 Watts each)</td>
<td>0.1</td>
<td>340</td>
</tr>
<tr>
<td></td>
<td>Peripheral Media Tower (PMT)</td>
<td>0.13</td>
<td>425</td>
</tr>
<tr>
<td></td>
<td>TOTAL</td>
<td>1</td>
<td>3625</td>
</tr>
</tbody>
</table>

*Technical Room is not mandatory, the placements of these elements are recommended in the Exam Room.

**WARNING**

This chart contains only the principal components of the PET/CT system and does not include information about non-GE supplied equipment.
CONNECTIVITY REQUIREMENTS

Broadband Connections are necessary between customer’s imaging devices and the GE Support Center, starting from the installation process to ensure full support from the Engineering Teams. GE provides remote maintenance and maximum availability for the customer’s system, during the equipment’s full lifetime. GE guarantees to keep the equipment at a maximum performance level.

Proactive and reactive maintenance are available through utilizing a wide range of digital tools. You may choose from the connectivity solutions listed below:

- Site-to-Site VPN/GE Solution
- Site-to-Site VPN/Customer Solution
- Connection through Dedicated Service Network
- Internet Access - connectivity for InSite 2.0

The requirements for these connectivity solutions are explained in the broadband solutions catalogue (separate document).

ELECTRICAL NOTES

1. All wires specified shall be copper stranded, flexible, thermo-plastic, color coded, cut 10 foot long at outlet boxes, duct termination points or stubbed conduit ends. All conductors, power, signal and ground, must be run in a conduit or duct system. Electrical contractor shall ring out and tag all wires at both ends. Wire runs must be continuous copper stranded and free from splices.
   1.1. Aluminum or solid wires are not allowed.
   2. Wire sizes given are for use of equipment. Larger sizes may be required by local codes.
   3. It is recommended that all wires be color coded, as required in accordance with national and local electrical codes.
   4. Conduit sizes shall be verified by the architect, electrical engineer or contractor, in accordance with local or national codes.
   5. Convenience outlets are not illustrated. Their number and location are to be specified by others. Locate at least one convenience outlet close to the system control, the power distribution unit and one on each wall of the procedure room. Use hospital approved outlet or equivalent.
   6. General room illumination is not illustrated. Caution should be taken to avoid excessive heat from overhead spotlights. Damage can occur to ceiling mounting components and wiring if high wattage bulbs are used. Recommend low wattage bulbs no higher than 75 watts and use dimmer controls (except mr). Do not mount lights directly above areas where ceiling mounted accessories will be parked.
   7. Routing of cable ductwork, conduits, etc., must run direct as possible otherwise may result in the need for greater than standard cable lengths (refer to the interconnection diagram for maximum usable lengths point to point).
   8. Conduit turns to have large, sweeping bends with minimum radius in accordance with national and local electrical codes.
   9. A special grounding system is required in all procedure rooms by some national and local codes. It is recommended in areas where patients might be examined or treated under present, future, or emergency conditions. Consult the governing electrical code and confer with appropriate customer administrative personnel to determine the areas requiring this type of grounding system.
   10. The maximum point to point distances illustrated on this drawing must not be exceeded.
   11. Physical connection of primary power to GE equipment is to be made by customers electrical contractor with the supervision of a GE representative. The GE representative would be required to identify the physical connection location, and insure proper handling of GE equipment.
   12. GEHC conducts power audits to verify quality of power being delivered to the system. The customer’s electrical contractor is required to be available to support this activity.

- All junction boxes, conduit, duct, duct dividers, switches, circuit breakers, cable tray, etc., are to be supplied and installed by customers electrical contractor.
- Conduit and duct runs shall have sweep radius bends
- Conduits and duct above ceiling or below finished floor must be installed as near to ceiling or floor as possible to reduce run length.
- Ceiling mounted junction boxes illustrated on this plan must be installed flush with finished ceiling.
- All ductwork must meet the following requirements:
  1. Ductwork shall be metal with dividers and have removable, accessible covers.
  2. Ductwork shall be certified/rated for electrical power purposes.
  3. Ductwork shall be electrically and mechanically bonded together in an approved manner.
- PVC as a substitute must be used in accordance with all local and national codes.
- Electrical contractor to install pull cords for all cable run conduits between the equipment room and the operators control room.
- 10 foot pigtails at all junction points.
- Grounding is critical to equipment function and patient safety. Site must conform to wiring specifications shown on this plan.
### Additional Conduit Runs (Contractor Supplied and Installed)

<table>
<thead>
<tr>
<th>From</th>
<th>To</th>
<th>Qty</th>
<th>Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 phase power</td>
<td>Main disconnect</td>
<td>1</td>
<td>AS REQ'D</td>
</tr>
<tr>
<td>Main disconnect</td>
<td>Emergency off</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>Power Distribution Unit</td>
<td>Door Switch</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>Power Distribution Unit</td>
<td>Power Distribution Unit</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>Warning light</td>
<td>Warning light control</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>1 phase power</td>
<td>1 phase power</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>Injector</td>
<td>Injector Control</td>
<td>1</td>
<td>2 1/2</td>
</tr>
</tbody>
</table>

### Description of Items

1. Main disconnect panel (recommended 1 1/2m [60 in] floor to middle of panel)
2. 75 [3”] conduit below floor for water lines
3. 64 [2 1/2”] conduit below floor
4. 89 [3 1/2”] conduit below floor
5. 600 x 400 x 100 [12” x 16” x 4”] box for power distribution unit
6. 300 x 400 x 150 [12” x 16” x 6”] box for power distribution unit
7. 150 x 150 x 100 [6” x 6” x 4”] box for injector control
8. 150 x 150 x 100 [6” x 6” x 4”] box above ceiling for injector
9. 250 x 100 [10” x 3 1/2”] surface wall duct with minimum 2 dividers
10. 450 x 100 [18” x 3 1/2”] surface wall duct with minimum 2 dividers

### Outlet Legend for GE Equipment

- Dedicated telephone line(s)
- Network outlet
- Duplex hospital grade, dedicated wall outlet 120-v, single phase power
- System emergency off (SEO), (recommended height 1.2m [48"] above floor)
- X-Ray room warning light control panel
- X-Ray ON lamp (111) - 24V
- Door interlock switch (needed only if required by state/local codes)
The cables from signaling and remote control (Y, SEO, L...) will go to A1 Main Disconnect with a pigtail length of 1.5m, and will be connected during installation. Each conductor will be identified and isolated (screw connector).

GROUND SYSTEM
- System of equipotential grounding.
- Equipotential: The equipotential link will be by means of an equipotential bar. This equipotential bar should be connected to the protective earth conductors in the ducts of the non GE cableways and to additional equipotential connections linking up all the conducting units in the rooms where GE system units are located.
- The impedance of the earth bar should be less than or equal to 2 Ω (ohm).

CABLEWAYS
The general rules for laying cableways should meet the conditions laid down in current standards and regulations, with regard to:
- Protecting cables against water (cableways should be waterproof).
- Protecting cables against abnormal temperatures (proximity to heating pipes or ducts).
- Protecting cables against temperature shocks.
- Replacing cables (cableways should be large enough for cables to be replaced).
- Metal cableways should be grounded.

FEEDER TABLE
<table>
<thead>
<tr>
<th>MIN. FEEDER WIRE SIZE, AWG OR MCM (sq. mm)/VAC</th>
<th>MINIMUM FEEDER WIRE LENGTH - ft (m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 AWG (16)</td>
<td>150 (46)</td>
</tr>
<tr>
<td>8 AWG (25)</td>
<td>150 (46)</td>
</tr>
<tr>
<td>6 AWG (50)</td>
<td>150 (46)</td>
</tr>
</tbody>
</table>

In all cases qualified personnel must verify that the feeder (at the point of take-off) and the run to the CT-PET system meet all the requirements stated in the PIM.

For a single unit installation, the minimum transformer size is 100 kVA, with 3.4% rated regulation at unity power factor. Resultant maximum allowable feeder regulation is 3.4%.

Grounding conductor will be a 1/0 minimum. This ground will run from the equipment back to the power source/main grounding point and always travel in the same conduit with the feeders.