

Innova™ IGS 520, Innova™ IGS 530, Innova™ IGS 540

Pre-Installation Manual



OPERATING DOCUMENTATION

5499972-1-1EN
Revision 2

ATTENTION

LES APPAREILS A RAYONS X SONT DANGEREUX A LA FOIS POUR LE PATIENT ET POUR LE MANIPULATEUR SI LES MESURES DE PROTECTION NE SONT PAS STRICTEMENT APPLIQUEES

Bien que cet appareil soit construit selon les normes de sécurité les plus sévères, la source de rayonnement X représente un danger lorsque le manipulateur est non qualifié ou non averti.

Une exposition excessive au rayonnement X entraîne des dommages à l'organisme.

Par conséquent, toutes les précautions doivent être prises pour éviter que les personnes non autorisées ou non qualifiées utilisent cet appareil créant ainsi un danger pour les autres et pour elles-mêmes.

Avant chaque manipulation, les personnes qualifiées et autorisées à se servir de cet appareil doivent se renseigner sur les mesures de protection établies par la Commission Internationale de la Protection Radiologique, Annales 26 : Recommandations de la Commission Internationale sur la Protection Radiologique et les normes nationales en vigueur.

WARNING

X-RAY EQUIPMENT IS DANGEROUS TO BOTH PATIENT AND OPERATOR UNLESS MEASURES OF PROTECTION ARE STRICTLY OBSERVED

Though this equipment is built to the highest standards of electrical and mechanical safety, the useful x-ray beam becomes a source of danger in the hands of the unauthorized or unqualified operator.

Excessive exposure to x-radiation causes damage to human tissue.

Therefore, adequate precautions must be taken to prevent unauthorized or unqualified persons from operating this equipment or exposing themselves or others to its radiation.

Before operation, persons qualified and authorized to operate this equipment should be familiar with the Recommendations of the International Commission on Radiological Protection, contained in Annals Number 26 of the ICRP, and with applicable national standards.

ATENCION

LOS APARATOS DE RAYOS X SON PELIGROSOS PARA EL PACIENTE Y EL MANIPULADOR CUANDO LAS NORMAS DE PROTECCION NO ESTAN OBSERVADAS

Aunque este aparato está construido según las normas de seguridad más estrictas, la radiación X constituye un peligro al ser manipulado por personas no autorizadas o incompetentes. Una exposición excesiva a la radiación X puede causar daños al organismo.

Por consiguiente, se deberán tomar todas las precauciones necesarias para evitar que las personas incompetentes o no autorizadas utilicen este aparato, lo que sería un peligro para los demás y para sí mismas.

Antes de efectuar las manipulaciones, las personas habilitadas y competentes en el uso de este aparato, deberán informarse sobre las normas de protección fijadas por la Comisión Internacional de la Protección Radiológica, Anales No 26: Recomendaciones de la Comisión Internacional sobre la Protección Radiológica y normas nacionales.

ACHTUNG

RÖNTGENAPPARATE SIND EINE GEFAHR FÜR PATIENTEN SOWIE BEDIENUNGSPERSONAL, WENN DIE GELTENDEN SICHERHEITSVORKEHRUNGEN NICHT GENAU BEACHTET WERDEN

Dieser Apparat entspricht in seiner Bauweise strengsten elektrischen und mechanischen Sicherheitsnormen, doch in den Händen unbefugter oder unqualifizierter Personen wird er zu einer Gefahrenquelle.

Übermäßige Röntgenbestrahlung ist für den menschlichen Organismus schädlich.

Deswegen sind hinreichende Vorsichtsmaßnahmen erforderlich, um zu verhindern, daß unbefugte oder unqualifizierte Personen solche Geräte bedienen oder sich selbst und andere Personen deren Bestrahlung aussetzen können.

Vor Inbetriebnahme dieses Apparats sollte sich das qualifizierte und befugte Bedienungspersonal mit den geltenden Kriterien für den gefahrlosen Strahleneinsatz durch sorgfältiges Studium des Hefts Nr. 26 der Internationalen Kommission für Strahlenschutz (ICRP) vertraut machen: Empfehlungen der Internationalen Kommission für Strahlenschutz und anderer nationaler Normenbehörden.

Important Information

LANGUAGE

ПРЕДУПРЕЖДЕНИЕ (BG)	<p>Това упътване за работа е налично само на английски език.</p> <ul style="list-style-type: none">• Ако доставчикът на услугата на клиента изиска друг език, задължение на клиента е да осигури превод.• Не използвайте оборудването, преди да сте се консултирали и разбрали упътването за работа.• Неспазването на това предупреждение може да доведе до нараняване на доставчика на услугата, оператора или пациента в резултат на токов удар, механична или друга опасност.
警告 (ZH-CN)	<p>本维修手册仅提供英文版本。</p> <ul style="list-style-type: none">• 如果客户的维修服务人员需要非英文版本，则客户需自行提供翻译服务。• 未详细阅读和完全理解本维修手册之前，不得进行维修。• 忽略本警告可能对维修服务人员、操作人员或患者造成电击、机械伤害或其他形式的伤害。
警告 (ZH-HK)	<p>本服務手冊僅提供英文版本。</p> <ul style="list-style-type: none">• 倘若客戶的服務供應商需要英文以外之服務手冊，客戶有責任提供翻譯服務。• 除非已參閱本服務手冊及明白其內容，否則切勿嘗試維修設備。• 不遵從本警告或會令服務供應商、網絡供應商或病人受到觸電、機械性或其他的危險。
警告 (ZH-TW)	<p>本維修手冊僅有英文版。</p> <ul style="list-style-type: none">• 若客戶的維修廠商需要英文版以外的語言，應由客戶自行提供翻譯服務。• 請勿試圖維修本設備，除非 您已查閱並瞭解本維修手冊。• 若未留意本警告，可能導致維修廠商、操作員或病患因觸電、機械或其他危險而受傷。
UPOZORENJE (HR)	<p>Ovaj servisni priručnik dostupan je na engleskom jeziku.</p> <ul style="list-style-type: none">• Ako davatelj usluge klijenta treba neki drugi jezik, klijent je dužan osigurati prijevod.• Ne pokušavajte servisirati opremu ako niste u potpunosti pročitali i razumjeli ovaj servisni priručnik.• Zanemarite li ovo upozorenje, može doći do ozljede davatelja usluge, operatera ili pacijenta uslijed strujnog udara, mehaničkih ili drugih rizika.

VÝSTRAHA (CS)	Tento provozní návod existuje pouze v anglickém jazyce. <ul style="list-style-type: none">• V případě, že externí služba zákazníkům potřebuje návod v jiném jazyce, je zajištěný překlad do odpovídajícího jazyka úkolem zákazníka.• Nesnažte se o údržbu tohoto zařízení, aniž byste si přečetli tento provozní návod a pochopili jeho obsah.• V případě nedodržování této výstrahy může dojít k poranění pracovníka prodejního servisu, obslužného personálu nebo pacientů vlivem elektrického proudu, respektive vlivem mechanických či jiných rizik.
ADVARSEL (DA)	Denne servicemanual findes kun på engelsk. <ul style="list-style-type: none">• Hvis en kundes tekniker har brug for et andet sprog end engelsk, er det kundens ansvar at sørge for oversættelse.• Forsøg ikke at servicere udstyret uden at læse og forstå denne servicemanual.• Manglende overholdelse af denne advarsel kan medføre skade på grund af elektrisk stød, mekanisk eller anden fare for teknikeren, operatøren eller patienten.
WAARSCHUWING (NL)	Deze onderhoudshandleiding is enkel in het Engels verkrijgbaar. <ul style="list-style-type: none">• Als het onderhoudspersoneel een andere taal vereist, dan is de klant verantwoordelijk voor de vertaling ervan.• Probeer de apparatuur niet te onderhouden alvorens deze onderhoudshandleiding werd geraadpleegd en begrepen is.• Indien deze waarschuwing niet wordt opgevolgd, zou het onderhoudspersoneel, de operator of een patiënt gewond kunnen raken als gevolg van een elektrische schok, mechanische of andere gevaren.
WARNING (EN)	This service manual is available in English only. <ul style="list-style-type: none">• If a customer's service provider requires a language other than English, it is the customer's responsibility to provide translation services.• Do not attempt to service the equipment unless this service manual has been consulted and is understood.• Failure to heed this warning may result in injury to the service provider, operator or patient from electric shock, mechanical or other hazards.
HOIATUS (ET)	See teenindusjuhend on saadaval ainult inglise keeles. <ul style="list-style-type: none">• Kui klienditeeninduse osutaja nõub juhendit inglise keelest erinevas keeles, vastutab klient tõlketeenuse osutamise eest.• Ärge üritage seadmeid teenindada enne eelnevalt käesoleva teenindusjuhendiga tutvumist ja sellest aru saamist.• Käesoleva hoiatuse eiramise võib põhjustada teenuseosutaja, operaatori või patiendi vigastamist elektrilöögi, mehaanilise või muu ohu tagajärvel.

VAROITUS (FI)	Tämä huolto-ohje on saatavilla vain englanniksi. <ul style="list-style-type: none">• Jos asiakkaan huoltohenkilöstö vaatii muuta kuin englanninkielistä materiaalia, tarvittavan käänökseen hankkiminen on asiakkaan vastuulla.• Älä yritä korjata laitteistoa ennen kuin olet varmasti lukenut ja ymmärtänyt tämän huolto-ohjeen.• Mikäli tästä varoitusta ei noudateta, seurauksena voi olla huoltohenkilöstön, laitteiston käyttäjän tai potilaan vahingoittuminen sähköiskun, mekaanisen vian tai muun vaaratilanteen vuoksi.
ATTENTION (FR)	Ce manuel d'installation et de maintenance est disponible uniquement en anglais. <ul style="list-style-type: none">• Si le technicien d'un client a besoin de ce manuel dans une langue autre que l'anglais, il incombe au client de le faire traduire.• Ne pas tenter d'intervenir sur les équipements tant que ce manuel d'installation et de maintenance n'a pas été consulté et compris.• Le non-respect de cet avertissement peut entraîner chez le technicien, l'opérateur ou le patient des blessures dues à des dangers électriques, mécaniques ou autres.
WARNUNG (DE)	Diese Serviceanleitung existiert nur in englischer Sprache. <ul style="list-style-type: none">• Falls ein fremder Kundendienst eine andere Sprache benötigt, ist es Aufgabe des Kunden für eine entsprechende Übersetzung zu sorgen.• Versuchen Sie nicht diese Anlage zu warten, ohne diese Serviceanleitung gelesen und verstanden zu haben.• Wird diese Warnung nicht beachtet, so kann es zu Verletzungen des Kundendiensttechnikers, des Bedieners oder des Patienten durch Stromschläge, mechanische oder sonstige Gefahren kommen.
ΠΡΟΕΙΔΟΠΟΙΗΣΗ (EL)	To παρόν εγχειρίδιο σέρβις διατίθεται μόνο στα αγγλικά. <ul style="list-style-type: none">• Εάν ο τεχνικός σέρβις ενός πελάτη απαιτεί το παρόν εγχειρίδιο σε γλώσσα εκτός των αγγλικών, αποτελεί ευθύνη του πελάτη να παρέχει τις υπηρεσίες μετάφρασης.• Μην επιχειρήσετε την εκτέλεση εργασιών σέρβις στον εξοπλισμό αν δεν έχετε συμβουλευτεί και κατανοήσει το παρόν εγχειρίδιο σέρβις.• Αν δεν προσέξετε την προειδοποίηση αυτή, ενδέχεται να προκληθεί τραυματισμός στον τεχνικό σέρβις, στο χειριστή ή στον ασθενή από ηλεκτροπληξία, μηχανικούς ή άλλους κινδύνους.
FIGYELMEZTETÉS (HU)	Ezen karbantartási kézikönyv kizárolag angol nyelven érhető el. <ul style="list-style-type: none">• Ha a vevő szolgáltatója angoltól eltérő nyelvre tart igényt, akkor a vevő felelőssége a fordítás elkészítése.• Ne próbálja elkezdeni használni a berendezést, amíg a karbantartási kézikönyvben leírtakat nem értelmeztek.• Ezen figyelmeztetés figyelmen kívül hagyása a szolgáltató, működtető vagy a beteg áramütés, mechanikai vagy egyéb veszélyhelyzet miatti sérülését eredményezheti.

AÐVÖRUN (IS)	<p>Þessi þjónustuhandbók er aðeins fáanleg á ensku.</p> <ul style="list-style-type: none">Ef að þjónustuveitandi viðskiptamanns þarfnast annas tungumáls en ensku, er það skylda viðskiptamanns að skaffa tungumálapjónustu.Reynið ekki að afgreiða tækið nema að þessi þjónustuhandbók hefur verið skoðuð og skilin.Brot á sinna þessari aðvörun getur leitt til meiðsla á þjónustuveitanda, stjórnanda eða sjúklings frá raflosti, vélrænu eða öðrum áhættum.
AVVERTENZA (IT)	<p>Il presente manuale di manutenzione è disponibile soltanto in lingua inglese.</p> <ul style="list-style-type: none">Se un addetto alla manutenzione richiede il manuale in una lingua diversa, il cliente è tenuto a provvedere direttamente alla traduzione.Procedere alla manutenzione dell'apparecchiatura solo dopo aver consultato il presente manuale ed averne compreso il contenuto.Il mancato rispetto della presente avvertenza potrebbe causare lesioni all'addetto alla manutenzione, all'operatore o ai pazienti provocate da scosse elettriche, urti meccanici o altri rischi.
警告 (JA)	<p>このサービスマニュアルには英語版しかありません。</p> <ul style="list-style-type: none">サービスを担当される業者が英語以外の言語を要求される場合、翻訳作業はその業者の責任で行うものとさせていただきます。このサービスマニュアルを熟読し理解せずに、装置のサービスを行わないでください。この警告に従わない場合、サービスを担当される方、操作員あるいは患者さんが、感電や機械的又はその他の危険により負傷する可能性があります。
경고 (KO)	<p>본 서비스 매뉴얼은 영어로만 이용하실 수 있습니다.</p> <ul style="list-style-type: none">고객의 서비스 제공자가 영어 이외의 언어를 요구할 경우, 번역 서비스를 제공하는 것은 고객의 책임입니다.본 서비스 매뉴얼을 참조하여 숙지하지 않은 이상 해당 장비를 수리하려고 시도하지 마십시오.본 경고 사항에 유의하지 않으면 전기 쇼크, 기계적 위험, 또는 기타 위험으로 인해 서비스 제공자, 사용자 또는 환자에게 부상을 입힐 수 있습니다.
BRĪDINĀJUMS (LV)	<p>Šī apkopes rokasgrāmata ir pieejama tikai angļu valodā.</p> <ul style="list-style-type: none">Ja klienta apkopes sniedzējam nepieciešama informācija citā valodā, klienta pienākums ir nodrošināt tulkojumu.Neveiciet aprīkojuma apkopi bez apkopes rokasgrāmatas izlasīšanas un saprašanas.Šī brīdinājuma neievērošanas rezultātā var rasties elektriskās strāvas trieciena, mehānisku vai citu faktoru izraisītu traumu risks apkopes sniedzējam, operatoram vai pacientam.

 ISPĖJIMAS (LT)	<p>Šis eksplotavimo vadovas yra tik anglų kalba.</p> <ul style="list-style-type: none">• Jei kliento paslaugų tiekėjas reikalauja vadovo kita kalba – ne anglų, suteikti vertimo paslaugas privalo klientas.• Neméginkite atlkti įrangos techninės priežiūros, jei neperskaitėte ar nesupratote šio eksplotavimo vadovo.• Jei nepaisysite šio įspėjimo, galimi paslaugų tiekėjo, operatoriaus ar paciento sužalojimai dėl elektros šoko, mechaninių ar kitų pavojų.
ADVARSEL (NO)	<p>Denne servicehåndboken finnes bare på engelsk.</p> <ul style="list-style-type: none">• Hvis kundens serviceleverandør har bruk for et annet språk, er det kundens ansvar å sørge for oversettelse.• Ikke forsøk å reparere utstyret uten at denne servicehåndboken er lest og forstått.• Manglende hensyn til denne advarselen kan føre til at serviceleverandøren, operatøren eller pasienten skades på grunn av elektrisk støt, mekaniske eller andre farer.
OSTRZEŻENIE (PL)	<p>Niniejszy podręcznik serwisowy dostępny jest jedynie w języku angielskim.</p> <ul style="list-style-type: none">• Jeśli serwisant klienta wymaga języka innego niż angielski, zapewnienie usługi tłumaczenia jest obowiązkiem klienta.• Nie próbować serwisować urządzenia bez zapoznania się z niniejszym podręcznikiem serwisowym i zrozumienia go.• Niezastosowanie się do tego ostrzeżenia może doprowadzić do obrażeń serwisa, operatora lub pacjenta w wyniku porażenia prądem elektrycznym, zagrożenia mechanicznego bądź innego.
ATENÇÃO (PT-BR)	<p>Este manual de assistência técnica encontra-se disponível unicamente em inglês.</p> <ul style="list-style-type: none">• Se outro serviço de assistência técnica solicitar a tradução deste manual, caberá ao cliente fornecer os serviços de tradução.• Não tente reparar o equipamento sem ter consultado e compreendido este manual de assistência técnica.• A não observância deste aviso pode ocasionar ferimentos no técnico, operador ou paciente decorrentes de choques elétricos, mecânicos ou outros.
ATENÇÃO (PT-PT)	<p>Este manual de assistência técnica só se encontra disponível em inglês.</p> <ul style="list-style-type: none">• Se qualquer outro serviço de assistência técnica solicitar este manual noutra idioma, é da responsabilidade do cliente fornecer os serviços de tradução.• Não tente reparar o equipamento sem ter consultado e compreendido este manual de assistência técnica.• O não cumprimento deste aviso pode colocar em perigo a segurança do técnico, do operador ou do paciente devido a choques eléctricos, mecânicos ou outros.

ATENȚIE (RO)	<p>Acest manual de service este disponibil doar în limba engleză.</p> <ul style="list-style-type: none">• Dacă un furnizor de servicii pentru clienți necesită o altă limbă decât cea engleză, este de datoria clientului să furnizeze o traducere.• Nu încercați să reparați echipamentul decât ulterior consultării și înțelegerea acestui manual de service.• Ignorarea acestui avertisment ar putea duce la rănirea depanatorului, operatorului sau pacientului în urma pericolelor de electrocutare, mecanice sau de altă natură.
ОСТОРОЖНО! (RU)	<p>Данное руководство по техническому обслуживанию представлено только на английском языке.</p> <ul style="list-style-type: none">• Если сервисному персоналу клиента необходимо руководство не на английском, а на каком-то другом языке, клиенту следует самостоятельно обеспечить перевод.• Перед техническим обслуживанием оборудования обязательно обратитесь к данному руководству и поймите изложенное в нем сведения.• Несоблюдение требований данного предупреждения может привести к тому, что специалист по техобслуживанию, оператор или пациент получит удар электрическим током, механическую травму или другое повреждение.
UPOZORENJE (SR)	<p>Ovo servisno uputstvo je dostupno samo na engleskom jeziku.</p> <ul style="list-style-type: none">• Ako klijentov serviser zahteva neki drugi jezik, klijent je dužan da obezbedi prevodičke usluge.• Ne pokušavajte da opravite uređaj ako niste pročitali i razumeli ovo servisno uputstvo.• Zanemarivanje ovog upozorenja može dovesti do povređivanja servisera, rukovaoca ili pacijenta usled strujnog udara ili mehaničkih i drugih opasnosti.
UPOZORNENIE (SK)	<p>Tento návod na obsluhu je k dispozícii len v angličtine.</p> <ul style="list-style-type: none">• Ak zákazníkov poskytovateľ služieb vyžaduje iný jazyk ako angličtinu, poskytnutie prekladateľských služieb je zodpovednosťou zákazníka.• Nepokúšajte sa o obsluhu zariadenia, kým si neprečítate návod na obľahu a nepoznáte mu.• Zanedbanie tohto upozornenia môže spôsobiť zranenie poskytovateľa služieb, obsluhujúcej osoby alebo pacienta elektrickým prúdom, mechanické alebo iné ohrozenie.
ATENCION (ES)	<p>Este manual de servicio sólo existe en inglés.</p> <ul style="list-style-type: none">• Si el encargado de mantenimiento de un cliente necesita un idioma que no sea el inglés, el cliente deberá encargarse de la traducción del manual.• No se deberá dar servicio técnico al equipo, sin haber consultado y comprendido este manual de servicio.• La no observancia del presente aviso puede dar lugar a que el proveedor de servicios, el operador o el paciente sufran lesiones provocadas por causas eléctricas, mecánicas o de otra naturaleza.

VARNING (SV)	<p>Den här servicehandboken finns bara tillgänglig på engelska.</p> <ul style="list-style-type: none">Om en kunds servicetekniker har behov av ett annat språk än engelska, ansvarar kunden för att tillhandahålla översättningstjänster.Försök inte utföra service på utrustningen om du inte har läst och förstår den här servicehandboken.Om du inte tar hänsyn till den här varningen kan det resultera i skador på serviceteknikern, operatören eller patienten till följd av elektriska stötar, mekaniska faror eller andra faror.
OPOZORILO (SL)	<p>Ta servisni priročnik je na voljo samo v angleškem jeziku.</p> <ul style="list-style-type: none">Če ponudnik storitve stranke potrebuje priročnik v drugem jeziku, mora stranka zagotoviti prevod.Ne poskušajte servisirati opreme, če tega priročnika niste v celoti prebrali in razumeli.Če tega opozorila ne upoštevate, se lahko zaradi električnega udara, mehanskih ali drugih nevarnosti poškoduje ponudnik storitev, operater ali bolnik.
DİKKAT (TR)	<p>Bu servis kılavuzunun sadece ingilizcesi mevcuttur.</p> <ul style="list-style-type: none">Eğer müşteri teknisyeni bu kılavuzu ingilizce dışında bir başka lisandan talep ederse, bunu tercüme ettirmek müşteriye düşer.Servis kılavuzunu okuyup anlamadan ekipmanlara müdahale etmeyiniz.Bu uyarıya uyulmaması, elektrik, mekanik veya diğer tehlikelerden dolayı teknisyen, operatör veya hastanın yaralanmasına yol açabilir.

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Revision History

Part/Rev	Date	Reason for Change	Pages
5499972-1-1EN rev 1	January 27, 2015	Initial release of direction 5499972-1-1EN	242
5499972-1-1EN rev 2	March 10, 2015	Second release of direction 5499972-1-1EN	244

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Chapter 1 General Requirements

1 Objectives & Overview

1.1 Object and Scope of this manual

1.1.1 Object and Scope

This document is intended as a guide and information resource to properly plan and prepare a site for the installation of an Innova system (This includes the Innova IGS 520, Innova IGS 530 and Innova IGS 540).

In addition, this document provides references to the pre-installation documents of the various product included with an Innova System.

These documents are intended to assist the Installation Specialist and the Site Planner in properly preparing a site for the installation of this system.

It provides pre-installation data, such as site preparation prior to the delivery of the Innova System, environmental and electrical requirements and some additional planning aids.



WARNING

MAKE SURE THE ROOM PREPARATION COMPLIES WITH LOCAL REGULATIONS AS THE PIM IS NOT INTENDED TO REFLECT ALL OF THEM

1.1.2 Quebec

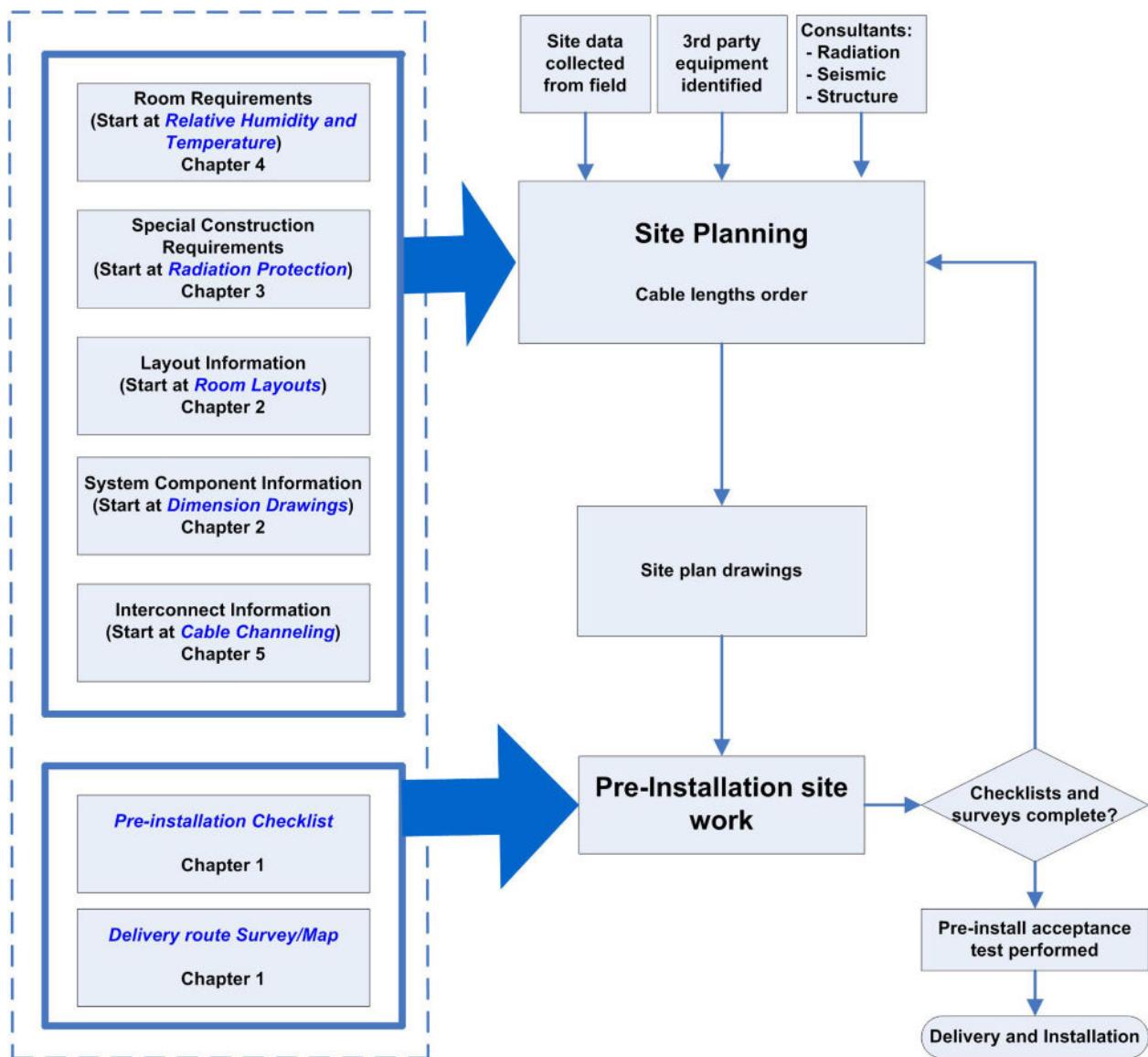
GE Healthcare is "GE Santé" in Province of Quebec - Canada.

1.2 Pre-Installation Process

Complete the checklists in ROOM LAYOUTS, ELECTRICAL REQUIREMENTS, and GENERAL REQUIREMENTS of this manual. They represent an important part of the pre-installation process. The checklists summarize the required preparations and allow to verify the proper completion of the pre-installation procedures.

You will find hereafter a chart of the information flow in the pre-installation process.

Illustration 1-1:



2 Customer Responsibilities

2.1 Responsibilities of the Purchaser/Customer

To ensure that the installation of an Innova System meets the purchaser or customer expectations, it is important to determine who will take responsibility for the various items during the system installation process. To help you in determining these responsibilities, review the following checklists with the customer and assign responsibilities as appropriate:

- Tool and Test Equipment ([Tools and Test Equipment](#))
- Pre-Installation Checklist ([Pre-Installation Checklist](#))

Contract Changes:

Be sure to inform the customer that the cost of any alteration or modification not specified in the sales contract are liable to the customer.

The following GE-supplied equipment must be installed by the Hospital's Contractors, per room drawings:

- PDB (Electrical Power Distribution Box or Main Disconnect Panel)
- Gantry & Table baseplate hole drilling (Per supplied template)
- LCD Monitor suspension stationary rails
- Gantry baseplate grout
- Gantry baseplate
- Omega/Tilting table common baseplate (if applicable)

NOTE: In case of Tilting table upgrade, it is critical to have the table baseplate flushed in the concrete.

2.2 Equipment Classifications

The following equipment classifications are applicable to the product:

Classification category	Equipment classification
Protection against electric shock	Class I   TO AVOID THE RISK OF ELECTRIC SHOCK, THIS EQUIPMENT MUST ONLY BE CONNECTED TO A SUPPLY MAINS WITH PROTECTIVE EARTH.
Degree of protection against electric shock	 Type B applied parts Applied parts complying with the specified requirements of the IEC 60601-1 standard to provide protection against electric shock, particularly regarding allowable patient leakage current and patient auxiliary current include: <ul style="list-style-type: none">• Mattress• Table accessories: shoulder rest, foot rest, table head extender, Removable rails (sleeve), Head widener with pad/cushion, Width extender with pad/cushion, armboard with thick pad/cushion, rail extender and patient restraint strap with cushion.
Degree of protection against harmful ingress of water	Ordinary equipment (enclosed equipment without protection against ingress of water); except footswitch which is a watertight device (protected against the effects of submersion, IPX7/IPX8) Systems with OR Table: Table, TSSC, Smart Box, Table Panning Device and Innova Central touch screen are watertight devices protected against the effects of splashing (IPX4). Footswitch is a watertight device protected against the effect of submersion (IPX8).
Method(s) of sterilization or disinfection recommended by the manufacturer	<ul style="list-style-type: none">• Sterilization: not applicable• Disinfection: refer to Operator Manual (Chapter Safety and Regulatory, section Disinfection), Recommended disinfecting agents. The system does not fulfill the requirements for AP/APG classification (IEC 60601-1).
Degree of safety of application in the presence of a flammable anesthetic mixture with air or with oxygen or nitrous oxide	Equipment not suitable for use in the presence of a flammable anesthetic mixture with air or with oxygen or nitrous oxide
Mode of operation	Continuous operation with intermittent loading

The VIVID E9 ultrasound unit is a Class I device, type CF, according to Sub-clause 14 of IEC60601-1 (1988).

For further details please refer *VIVID E9 Service Manual GA091568TPH* Table 1-17



NOTICE

The system can only be installed in an anesthetizing location if that location is classified as Other Than Hazardous as per NFPA 70 clause 517.60



NOTICE

The product is not classified as AP, APG (Equipment not suitable for use in the presence of a flammable anaesthetic mixture with air or with oxygen or nitrous oxide).

2.3 Pre-Installation Checklist

GE Healthcare Site Readiness Checklist							
GEHC Global Order # :		Customer:					
GEHC On-site Representative :		MI Supplier:					
Name of customer reviewed with :		Lead Installer:					
GEHC PMI :		Phone Number:					
Target Site Prep Completion Date:		Helper:					
The customer is responsible for proper site preparation and site readiness regardless of any GEHC inspections/assessments. It is under its responsibility to ensure electrical installation is compliant to local regulations.							
For MR Magnet Delivery: Ensure cryogen vents, power for the cooling system and exhaust fan system are installed and operational (0.7T, 1.5T & 3T) and chilled water supply is available 24x7 that meets system cooling equipment requirements.							
Inspection Date:							
Item #	GEHC Minimum Requirements	Storage: Is item ready?	Predict (Pre-ship)		Verify (Delivery): Is item ready?	Validate (Mech Install): Is item ready?	Comments If "N", please enter in comments or action plan
			Is this item ready?	Will item be ready?			
1	Equipment installation drawings must match actual room size and must meet clearance requirements. Deviations that meet installation requirements may be red-lined, if red-lining is allowed by local code. Seismic requirements are identified on construction drawings.	X					
2	Delivery route to installation or storage area meets requirements and has been discussed and scheduled with the customer. Ensure floor protection is discussed, requirements identified, and will be available at time of delivery and installation.					X	
3	Rooms that will contain equipment, including storage areas, are dust free. Room security to prevent unauthorized access and theft has been discussed with customer. The customer is aware of these security issues, implications and responsibility.						
4	In room HVAC ductwork and units (in room) must be mechanically installed and dust free. Installation rooms appear to meet environmental conditions (see Further Definitions) and observed issues have been communicated to the customer. If being stored, storage area must meet PIM storage criteria.						
5	Ceiling grid is installed, Unistrut is located per the installation drawings, and permanent lighting is installed and operational.	X					

6	Floor is clean and prepared for final floor covering. Customer has verified floor leveling meets the equipment installation drawings and PIM specs and no visible defects are observed. Gantry and table baseplate are installed prior to delivery (if applicable)	X					
7	Access to a working phone at the facility for emergency use, including MR magnet delivery.	X					
8	All walls primed (final coat not needed on Day 1), and counter tops that will support equipment must be installed. No dust-producing cabinetry work in installation areas.	X					
9	Mechanical supplier has been provided with a set of equipment installation drawings for reference. For California, permitted construction drawings or PMI-specified installation drawings are required.	X	X	X	X		
10	Conduit/electrical cable ducting/dividers/access flooring installed, with the exception of surface-mounted floor ducting. Wiring to the main disconnect panel is installed and compliant with equipment installation drawings or pre-installation manual.	X	X	X	X		
Issued Date: 7/9/07 Rev 11		GEHC Only: COE # (888) 799.7266 Option 5 (PMI Support)					

3 Delivery Requirements

3.1 Shipping Information

3.1.1 Product Shipping Information

Refer to the table below. To obtain shipping information for components not specified in the table, refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova System Compatability](#).

Table 1-1:

PRODUCT OR COMPO-NENT	DIMENSIONS MM (INCHES)			WEIGHT POUNDS (KILO-GRAMS)	METHOD OF SHIPMENT
	Height	Width	Depth		
Gantry	1950 (77)	2790 (110)	1155 (45.5)	2,340 (1060)	Shipping Dolly. See Illustration 1-2
	2300 (90.5)	2900 (114)	1380 (54.5)	2,645 (1200)	Air shipment. See Illustration 1-3
C1 Cabinet	2110 (83)	1080 (42.5)	820 (32.3)	1186 (538)	On pallet. See Illustration 1-4
C2 Cabinet	2110 (83)	1080 (42.5)	820 (32.3)	740.7 (336)	On pallet. See Illustration 1-4
Omega Table Base As-sembly	1240 (49)	960 (38)	2140 (84.2)	1,290 (585)	On pallet. See Illustration 1-5
Omega Table Top Assem-bly	220 (9)	3470 (137)	840 (33)	155 (70)	On pallet. See Illustration 1-5
Tilting table Base Assem-bly and covers	1160 (45.7)	1000 (39.4)	2150 (84.6)	1653 (750)	On pallet. See Illustration 1-7 and Illustration 1-8
DL User parts	1040 (41)	860 (33.9)	680 (26.8)	220 (100)	On pallet
Videostation VCR	290 (11.5)	540 (21.3)	410 (16.2)	210 (95)	On pallet
X-Ray tube housing	960 (37.7)	770 (30.3)	710 (28)	250 (113)	On pallet
Chiller 4100	1200 (47.2)	555 (21.8)	610 (24)	264.5 (120)	
Auto-tranformer (Coolix 4100 chiller)	340 (13.4)	370 (14.5)	304 (12)	97 (44)	
Gantry Requisites					On pallet
Cables					On pallet
Monitor susp. bridge	640 (25.2)	980 (38.6)	3060 (120.5)	445 (210)	On pallet
Monitor susp. rails	380 (15)	300 (12)	5960 (235)	355 (160)	On pallet
Large Display monitor	1050 (41.3)	1500 (59)	800 (31.4)	209 (95)	On pallet, see Illustration 1-9
Large Display cabinet	1600 (63)	950 (37.4)	750 (29.5)	423 (192)	On pallet, see Illustration 1-10
LD suspension	1100 (43.3)	1100 (43.3)	1850 (72.8)	860 (390)	On pallet
LD suspension handle	400 (15.7)	950 (37.4)	1650 (65)	15 (7)	Cardbord box
LD suspension 36m har-ness	230 (9)	800 (34.5)	800 (34.5)	134 (62)	On pallet
Fluoro UPS UL (*)	2100 (82.7)	890 (35)	1000 (39.4)	1235 (561)	On pallet

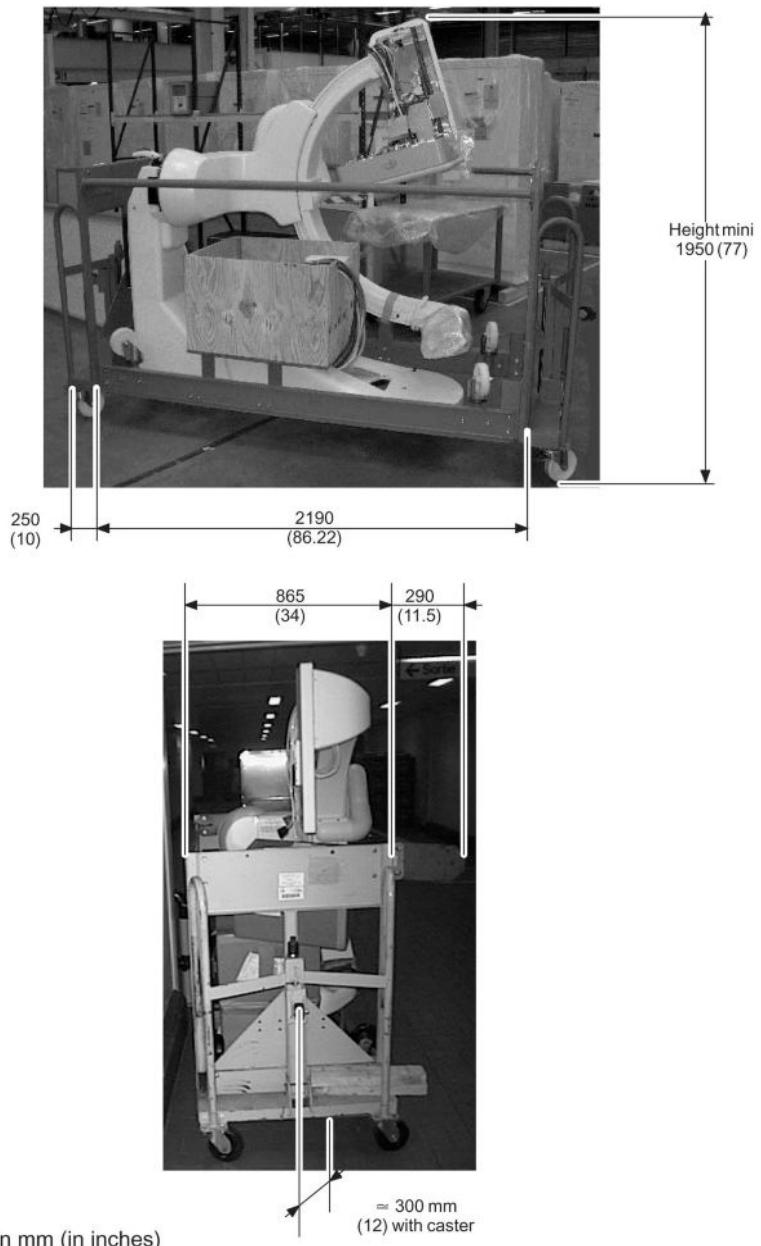
PRODUCT OR COMPO- NENT	DIMENSIONS MM (INCHES)			WEIGHT POUNDS (KILO- GRAMS)	METHOD OF SHIPMENT
	Height	Width	Depth		
Fluoro UPS CE (*)	1750 (68.9)	890 (35)	1000 (39.4)	1287 (585)	On pallet

(*) Estimated values

3.1.2 Detail Of Innova Shipping Information

3.1.2.1 Gantry On Shipping Dolly

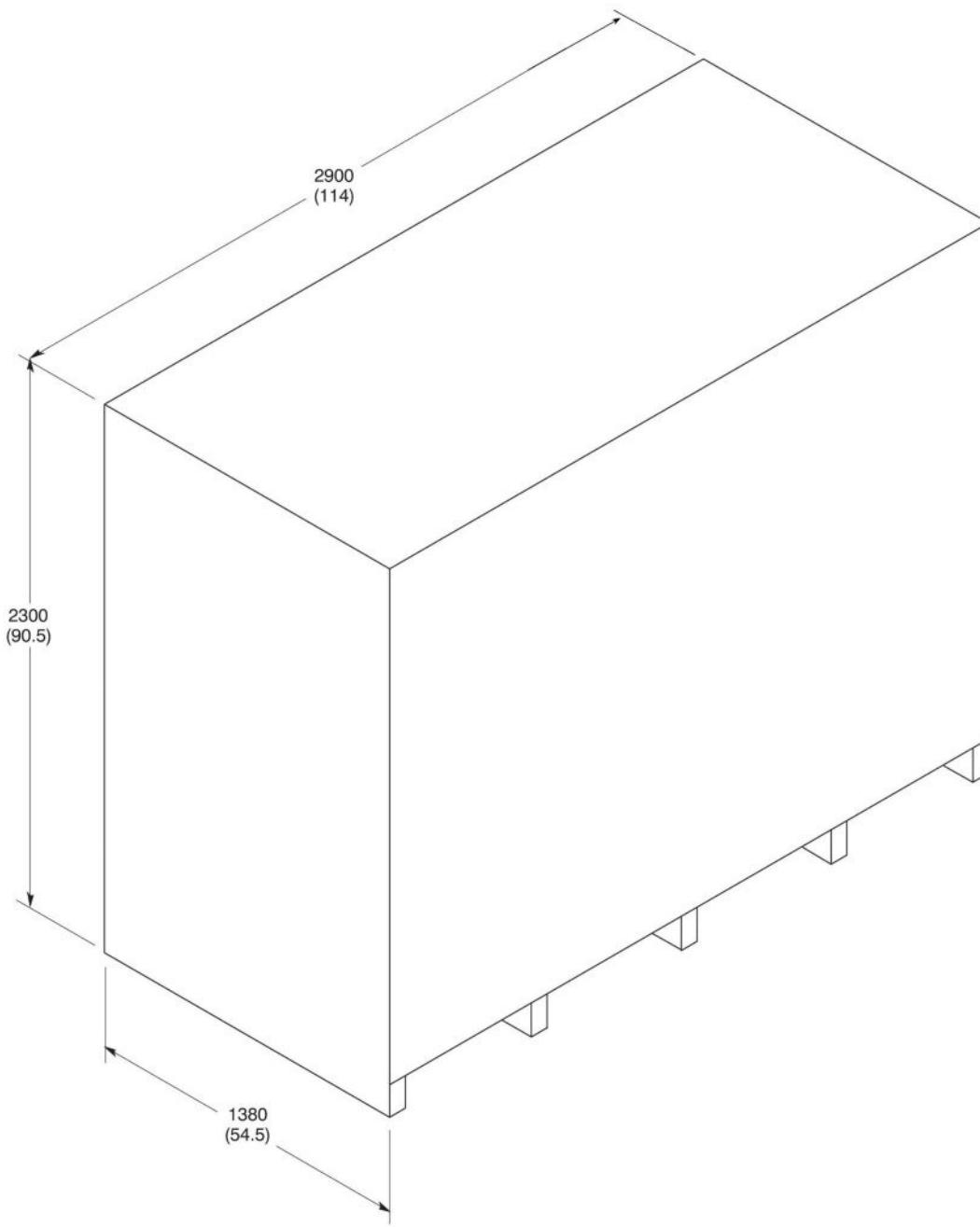
Illustration 1-2:



All dimensions are in mm (in inches)

3.1.2.2 Gantry Air Shipment

Illustration 1-3:



DIMENSIONS IN MM (INCHES)

NOT TO SCALE

3.1.2.3 C1 and C2 Cabinet in crate, on pallet

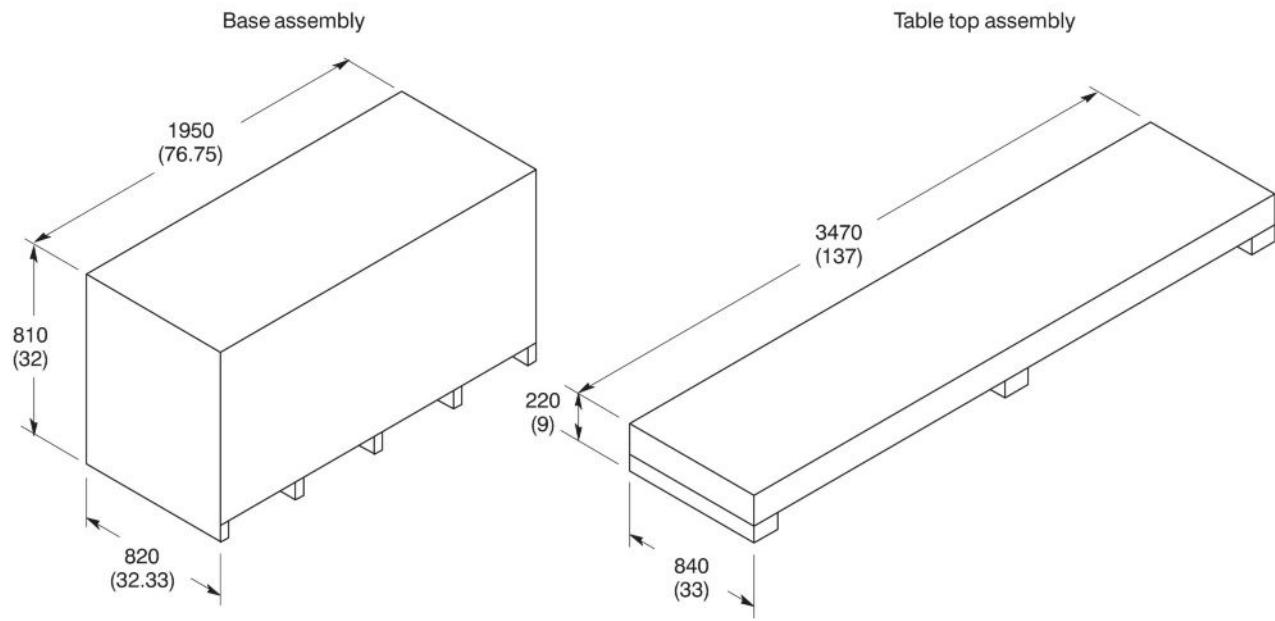
Illustration 1-4:



The shipping weight (crate and pallet) is 81kg (179 lbs).

3.1.2.4 Omega Shipment

Illustration 1-5:

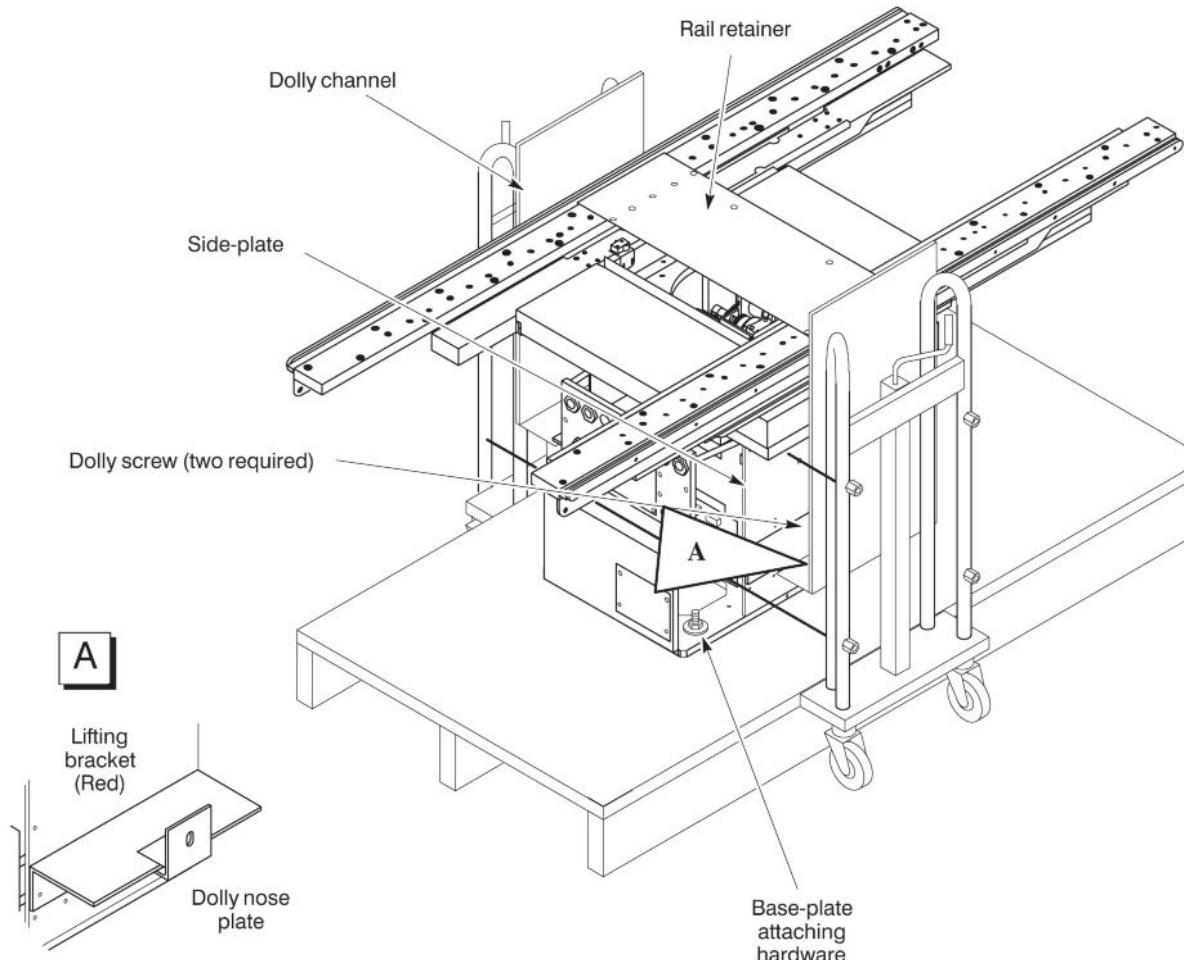


DIMENSIONS IN MM (INCHES)

NOT TO SCALE

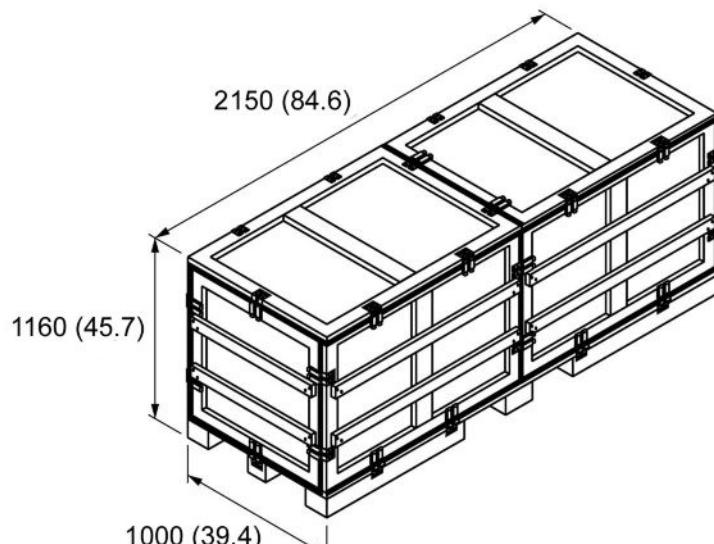
3.1.2.5 Omega Using Positioner Dollies

Illustration 1-6:



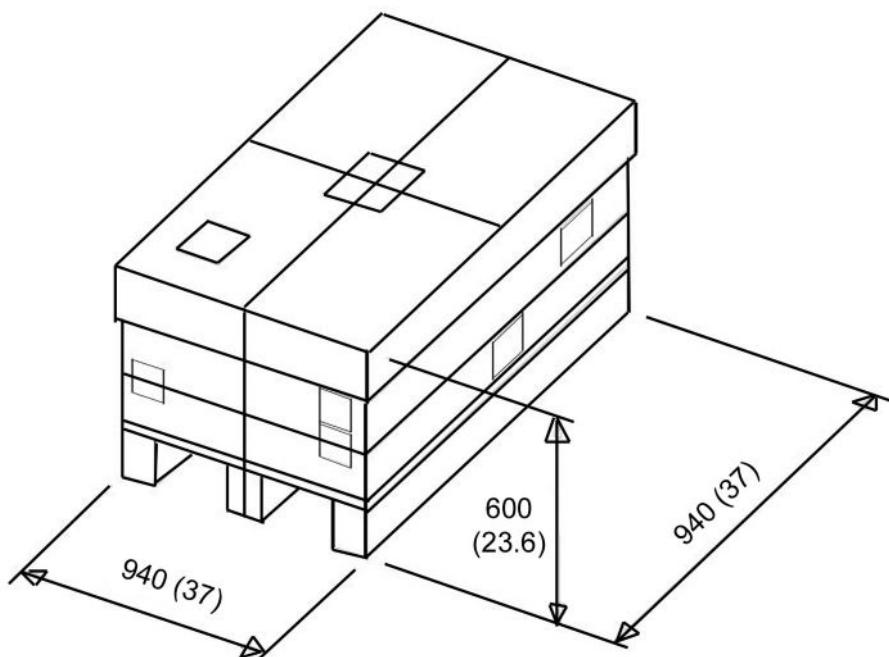
3.1.2.6 Innova^{IQ} Table and OR Table Shipment

Illustration 1-7: Table Shipment



All dimensions are in mm (inches)

Illustration 1-8: Covers Shipment



All dimensions are in mm (inches)

3.1.2.7 Large Display Option

Illustration 1-9: Large Display Monitor on pallet



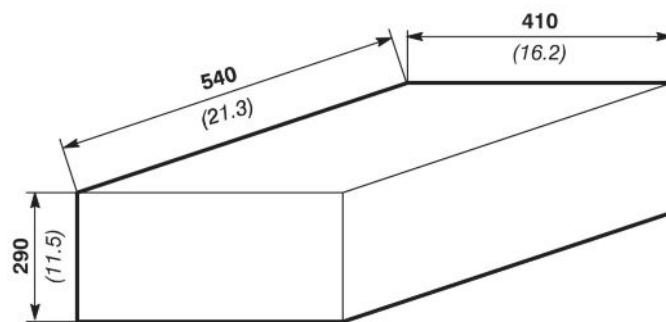
Illustration 1-10: Large Display cabinet on pallet



3.1.2.8 Other Elements Package

NOTE: All OEM parts are shipped inside their original boxes group as needed on pallets.

Illustration 1-11:



3.2 Tools and Test Equipment

Refer to [Table 1-2](#). To obtain a list of tools and test equipment for components not specified in [Table 1-2](#), refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova System Compatibility](#).

Table 1-2:

PRODUCT OR COM- PONENT	TOOL OR TEST EQUIP- MENT	USED FOR	SOURCE	RECEIVED (DATE)
Innova LC Positioner	Service Engineer's Tool Case	General Use		
	Level, Protractor Type	Positioner Checks		
	Plumb Line included in 46-216640G1	Positioner Checks		
	Torque Wrench 2 to 20 daN.m (15 ft. lbs. to 150 ft. lbs.)	Positioner Checks		
	1/2 inch Ratchet Wrench (2)	Raise and Lower Posi-tioner shipping dolly		
	Wrench, Spanner (46-176584P1)	High Voltage Cable In-stallation		
Status Display	Laptop Computer (MS-DOS Windows)	Positioner Configura-tion and Calibration		
	Same as for Innova LC (Service Engineer's Tool Case)			
Omega/Tilting table	Same as for Innova LC (Service Engineer's Tool Case) Fill in any additional tools or test equipment as required			
	Installation dolly (PN 5265134)	Replacing Omega with Tilting table		
C2 Cabinet	Same as for Innova LC (Service Engineer's Tool Case) Fill in any additional tools or test equipment as required			
X-Ray Head	Same as for Innova LC (Service Engineer's Tool Case) Fill in any additional tools or test equipment as required			
C1 Cabinet	Same as for Innova LC (Service Engineer's Tool Case) Fill in any additional tools or test equipment as required			
	Same as for Innova LC (Service Engineer's Tool Case) Fill in any additional tools or test equipment as required			
	Ethernet adaptation kit for laptop 2128794	General use (to be or-dered before delivery of system)		

PRODUCT OR COM- ONENT	TOOL OR TEST EQUIP- MENT	USED FOR	SOURCE	RECEIVED (DATE)
DL User parts				
Monitor Suspension				
Chiller	Phillips/Flathead screwdriver. Open chiller. Install wiring and hoses.			
Chiller Autotransformer (Coolix 4100)	Phillips/Flathead screwdriver. Open chiller autotransformer. Install wiring and hoses			
Large Display Lifting Tool (for Large Display Option)	Large Display Monitor Lifting Tool p/n 5418782	Raise Large Display Monitor for installation on Mavig suspension		

3.3 Door Size Requirements

Minimum door sizes also apply to hallways and elevators. For additional details, refer to [Shipping Information](#).

3.3.1 Door Height

The minimum door height (to accommodate Innova positioner on its dolly) is 1.980 m (78 in). If the height is limited to 2 m (79 in), you will need a fall over cabinet.

3.3.2 Door Width

The minimum door width needed (to accommodate the Innova LC shipping dolly) is:

- 1.165 m (46 in) with protective side rail,
- 1.096 m (43 in) with one protective side rail removed on site.

NOTE: Door widths are based on a *straight-in* approach requiring a 2.44 m (96 in) wide corridor. Calculations need to be made for accommodation of equipment through narrower corridors.

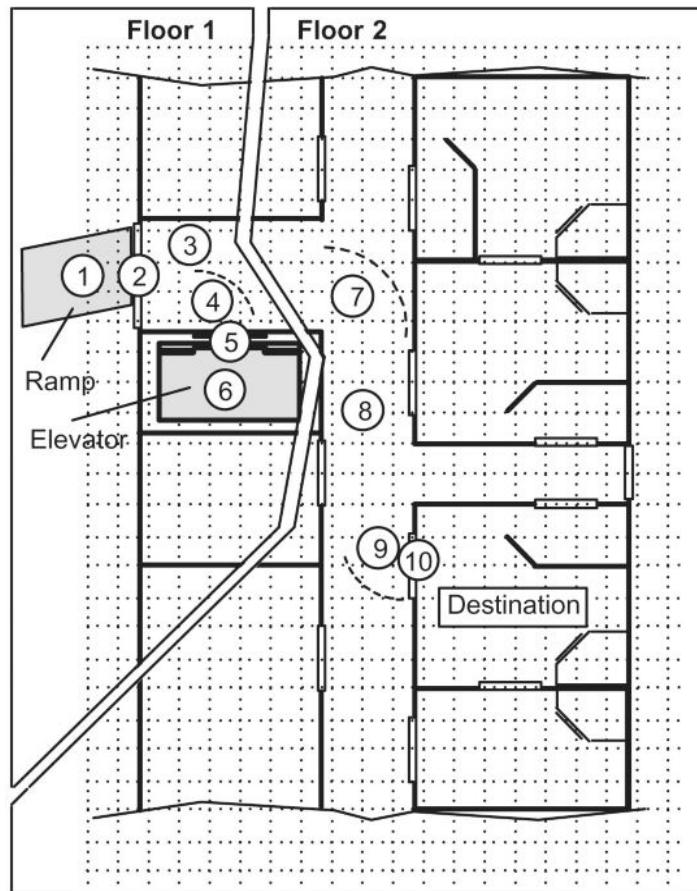
3.4 Route Survey

3.4.1 Step One — Sketch

Start preparing Route Survey by sketching a floor plan of the hospital or clinic which will receive the equipment. Include all areas on the delivery route from outside the building to destination. See [Illustration 1-12](#).

Reference Numbers: Numbers in circles refer to Route Survey data. The Route Survey is a form on which site data are listed (see [Section 3.4.2](#)).

Illustration 1-12:



3.4.2 Step Two — Survey

Data concerning the intended delivery route are recorded on the Route Survey in the following pages. Record all loading capacities, corridor widths, door openings, turning radii, flooring materials, elevator sizes, obstructions and so on.

3.4.3 Step Three — Check

Verify equipment can be transported via the route specified in [Section 3.4.1](#). Compare Route Survey compiled in [Section 3.4.2](#) to equipment specifications in this and other applicable pre-installation directions.

Table 1-3:

3.5 European Process Order Select

Several components like cables, stationary rails, etc. are length selectable. This selection should reflect the particularities encountered by an Installation Specialist on Site.

All cable and component selections must be sent at least 6 weeks before RSDD to your PMC. The selection tool is available on the website: [http://supportcentral.ge.com/products/
sup_products.asp?prod_id=99484](http://supportcentral.ge.com/products/sup_products.asp?prod_id=99484)

4 Product Storage and Handling Requirements

4.1 Product Storage and Handling limits



NOTICE

Avoid extremes in temperatures

To obtain Temperature, Humidity or Pressure requirements for components not specified in [Table 1-4](#), refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova LC System Compatibility](#).

Table 1-4: Storage and Transport

Component	TEMPERATURE		HUMIDITY (see Note (2))		PRESSURE	
	MIN	MAX	MIN	MAX	MIN	MAX
All UPS units (see also Section 4.2)	-20°C (-4°F)	+40°C (+104°F)	10%	90%	79.4 kPa	106 kPa
All Monitors	-20°C (-4°F)	+55°C (+131°F)	10%	80%	79.4 kPa	106 kPa
All Injectors	-20°C (-4°F)	+60°C (+140°F)	10%	80%	79.4 kPa	106 kPa
All Footswitches	-20°C (-4°F)	+60°C (+140°F)	10%	90%	79.4 kPa	106 kPa
Vivid E9 Option	-40°C (-40°F)	+70°C (+158°F)	30%	60%	79.4 kPa	106 kPa
Cardiac Collimator	-20°C (-4°F)	+70°C (+158°F)	10%	95%	79.4 kPa	106 kPa
Diamenter	-20°C (-4°F)	+60°C (+140°F)	0%	85%	79.4 kPa	106 kPa
All other components	-40°C (-40°F)	+70°C (+158°F)	10%	95%	79.4 kPa	106 kPa

NOTE: (1) The detector should be stored at 10 to 40 °C (50 to 104 °F) and less than or equal to 90% RH in the plastic wrapped shipping box. (This should include two bags of desiccant as well). The lowest temperature (e.g. 10 °C (50 °F)) and humidity is preferable. If they are to be stored outside of their shipping box or in the inner shipping box without plastic wrapping they should be stored at 20 °C (68 °F) or less and 30% RH or less. In terms of transportation, do not expose to temperatures below -20 °C (-4 °F) in its shipping box for more than 15 hours. The detector will reach the ambient temperature after 20 to 25 hours. The detector should not be allowed to reach temperatures less than -10 °C (14 °F) or irreparable damage to the detectors scintillator will occur. Care must be taken when removing a detector from a shipping box. If the detector has been subject to cold temperatures for an extended period the detector in the box should be allowed to sit in the plastic wrapped box to reach room temperature. This will prevent condensation from occurring. Condensation on the detector can cause irreparable damage to the electronics. Storage 10 to 40 °C (50 to 104 °F); 10 to 90 % RH, 250 day storage transportation -20 to +60 °C (-4 to 140 °F) and 10 to 80% RH. The Detector chiller is shipped within GEMS packaging.

NOTE: **(2) Special Humidity Instructions:** The following parts can be shipped in standard shipment conditions with the requirement that on arrival to installation site, and before supplying power to these parts, they shall be kept in an environmental relative humidity equal or lower than their specified capability, and that's for a minimum of 48 hours.

- 3kVA UPS 110 V & 220 V (Maximum specified relative humidity capability = 90%)
- LDM Monitor (Maximum specified relative humidity capability = 90%)
- CVI Injector (Maximum specified relative humidity capability = 85%)
- Dimentor (Maximum specified relative humidity capability = 80%)
- 1MP MX191 LCD Monitor (Maximum specified relative humidity capability = 80%)

4.2 Fluoro & 3KVA UPS Shipment Duration

NOTE: In the case the material shipped will be subject to different temperature values during shipment, the maximum shipment duration can be calculated, depending on the duration of different temperature values application, and using values given in the tables above

4.2.1 Fluoro UPS (CE and UL)

The maximum shipment duration is given in the table for both Fluoro UPS CE and UL. It depends on the temperature to which the material will be subject during shipment. This limitation is due to the capability of batteries included in UPS.

Table 1-5: Maximum shipment duration for Fluoro UPS

Temperature	Shipment max duration (Weeks)
55°C (131°F)	2
50°C (122°F)	3
40°C (104°F)	6
30°C (86°F)	12

4.2.2 3 KVA UPS (220 V and 110 V)

The maximum shipment duration is given in the table for both 3 KVA UPS 220 V and 110 V. It depends on the temperature to which the material will be subject during shipment. This limitation is due to the capability of batteries included in UPS.

Table 1-6: Maximum shipment duration for 3 KVA UPS

Temperature	Shipment max duration (Weeks)
55°C (131°F)	4
50°C (122°F)	7
40°C (104°F)	14
30°C (86°F)	25

4.3 Handling instructions

The packaging of the following components must be marked with special handling instructions for transport and storage.

- C1 Cabinet and C2 Cabinet:

Illustration 1-13:



- Gantry, Omega Table and Tilting Table:

Illustration 1-14:



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Chapter 2 Equipment Requirements

1 System Components

1.1 Presentation of the 3 Rooms

1.1.1 Examination room

- Gantry
- Omega / Tilting table
- Performix X-Ray Tube Assembly
- Collimator
- Innova Digital Detector



WARNING

THE ELECTRONIC CABINETS (C1, C2, OPTIONAL LD CABINET, COOLIX 4100 CHILLER, DETECTOR CONDITIONER AND FLUORO UPS WHEN INSTALLED) INCLUDE FANS THAT ARE CREATING AIR-CIRCULATION OF PULSED-AIR. WHEN THIS PULSED AIR IS IN AN ENVIRONMENT THAT MAY CONTAIN AIRBORNE PATHOGENS LIKE AN EXAM ROOM/CONTROL ROOM, THERE IS A RISK OF TRANSMISSION OF THESE AIRBORNE PATHOGENS FROM PATIENTS TO OTHER PATIENTS OR CLINICAL PERSONNEL (NOSOCOMIAL DISEASES).
TO REDUCE THIS RISK, THE ELECTRONIC CABINETS MUST BE INSTALLED IN A ROOM SEPARATED FROM EXAM ROOM/CONTROL ROOM, I.E., TECHNICAL ROOM.

1.1.2 Technical room

- C2 Cabinet
- C1 Cabinet
- Coolix 4100 chiller
- Detector Conditioner
- Fluoro UPS (option) (1 Cabinet)
- Fluoro UPS IF box (option)
- Large Display Cabinet and UPS (for Large Display Option)

1.1.3 Control room

- VCIM
- DL Flat Panel

- DL Keyboard
- Monitors

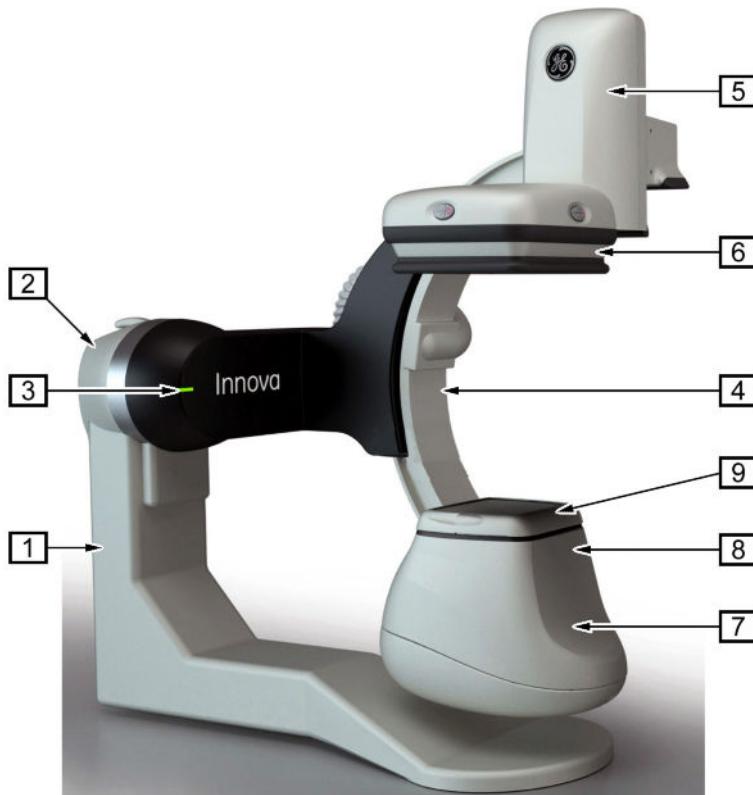
1.2 Description of Innova System

1.2.1 Description of the Gantry

A complete gantry comprises only three parts.

- C2 cabinet
- Gantry including an X-ray tube housing with a collimator and a Revolution Digital Detector.
- TSSC, Smart Box or Smart handle (options for remote installation).

Illustration 2-1:



- L-arm **1**,
- Pivot **2**,
- X-Ray LED **3**.
- C-arc **4**,
- Motorized elevator **5** for the Revolution Digital Detector,
- Revolution Digital Detector **6**:
 - 21 cm Revolution Digital Detector for Innova IGS 520.

- 31 cm Revolution Digital Detector for Innova IGS 530.
- 41 cm Revolution Digital Detector for Innova IGS 540.
- X-ray tube 
- Siemens collimator 
- X-Ray Tube cover spacer 

NOTE: Depending on country regulation (i.e. USA and New Zealand), the tube cover Spacer must be installed over the X-ray tube cover.

1.2.2 Description of the Patient table

1.2.2.1 Innova IGS 520 Case

Innova IGS 520 can be supplied with an OMEGA IV Compact patient table, an OMEGA V long patient table or a Tilting table.

1.2.2.2 Innova IGS 530 and Innova IGS 540 Case

Innova IGS 530 and Innova IGS 540 can be supplied with an OMEGA V Long patient table or Tilting table.

Illustration 2-2: Omega patient table (with TSSC)



Table system side Control (TSSC)

Illustration 2-3: Innova^{IQ} Table



TILT PATIENT TABLE

Illustration 2-4: OR Table



1.2.3 User Interfaces

Illustration 2-5: User interfaces with Omega table

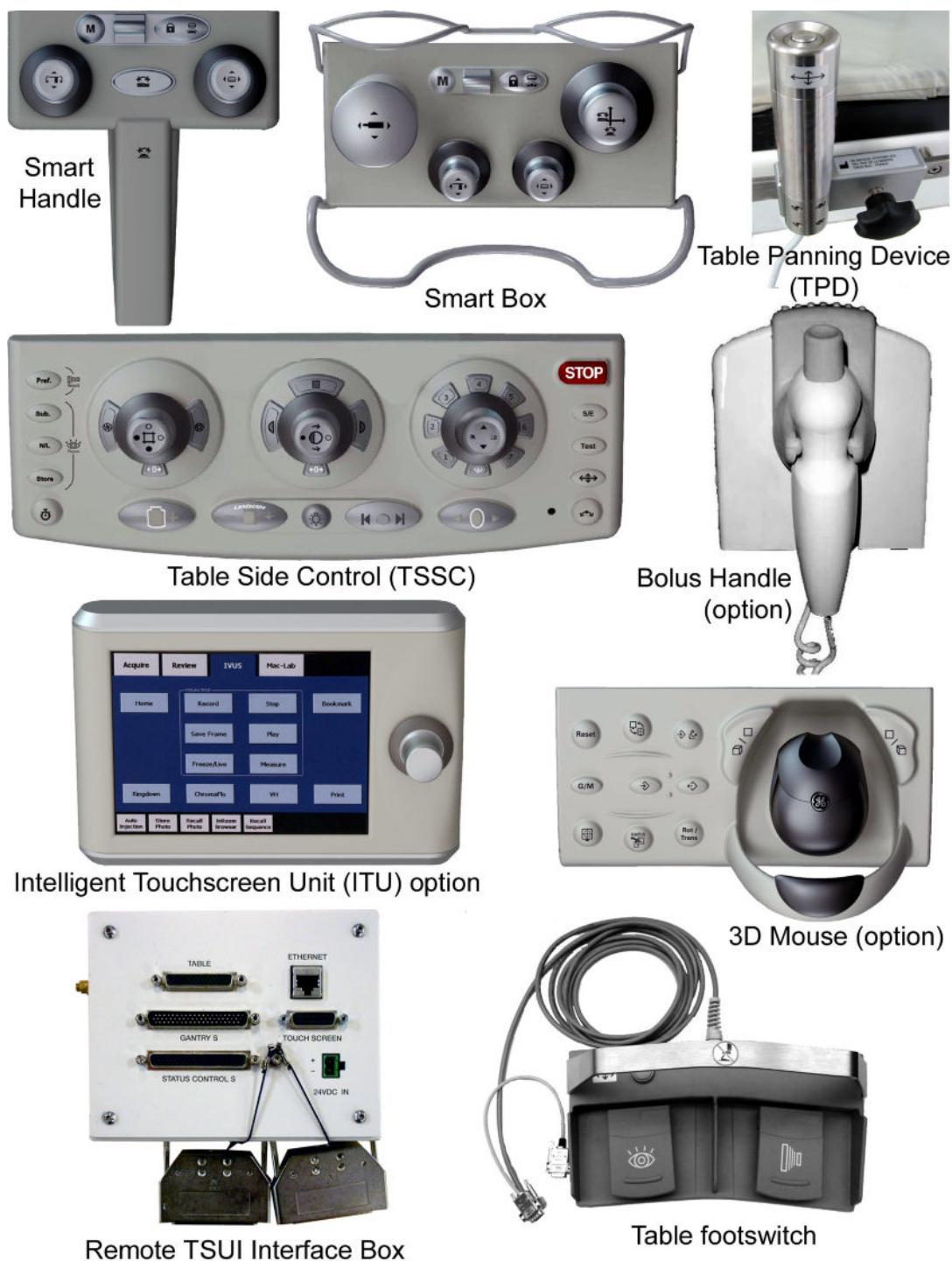


Illustration 2-6: User interfaces with Innova^{IQ} Table



Table Panning Device (TPD)



Table Side Control (TSSC)



Bolus Handle (option)



Intelligent Touchscreen Unit (ITU) option



3D Mouse (option)



Remote TSUI Interface Box



Table Footswitch

Illustration 2-7: User Interfaces with OR Table



Smart Box



Table Panning Device (TPD)



Table Side Control (TSSC)



Bolus Handle (option)



Intelligent Touchscreen Unit (ITU) option



3D Mouse (option)



Remote TSUI Interface Box



Table Footswitch

1.2.4 Accessories for Innova^{IQ} Table and OR Table

Illustration 2-8:



Illustration 2-9:



1.2.5 Description of the X-Ray generator

The Innova System uses a High Voltage System with component parts as follows according to the power requested (100 kW):

- Jedi - C1 cabinet,

- Ingrid H.V. tank mounted on the Performix 160A X-Ray tube housing.

1.2.6 Description of the X-Ray head

The Innova System uses a Performix 160A X-Ray tube housing:

- X-Ray tube
- collimator
- Ingrid HV tank
- oil/water exchanger
- contour filter
- tube chiller

NOTE: An external recirculating chiller is mandatory.

It is mandatory to place the Chiller in the Technical Room.

Illustration 2-10: Chiller Coolix 4100 (with auto-transformer)



1.2.7 Description of the Innova Imaging System

An Innova System is managed and controlled by a System including (does not list all components):

- C1 cabinet, inside which the main components are the RTAC, the DL, the Firewall, the HUB, the KVM, the 4 KVA PDU and the Jedi generator,
- C2 cabinet gantry and table control,
- Fluoro UPS interface box near C1 cabinet (Fluoro UPS option),
- VCIM console,
- Revolution Digital Detector mounted on Gantry:
- An external Digital Detector chiller is mandatory.

- LCD 19" monitors.
- LCD 58" EIZO GmbH LS580W Color Flat Panel (LDM option)

Illustration 2-11: C1 Cabinet

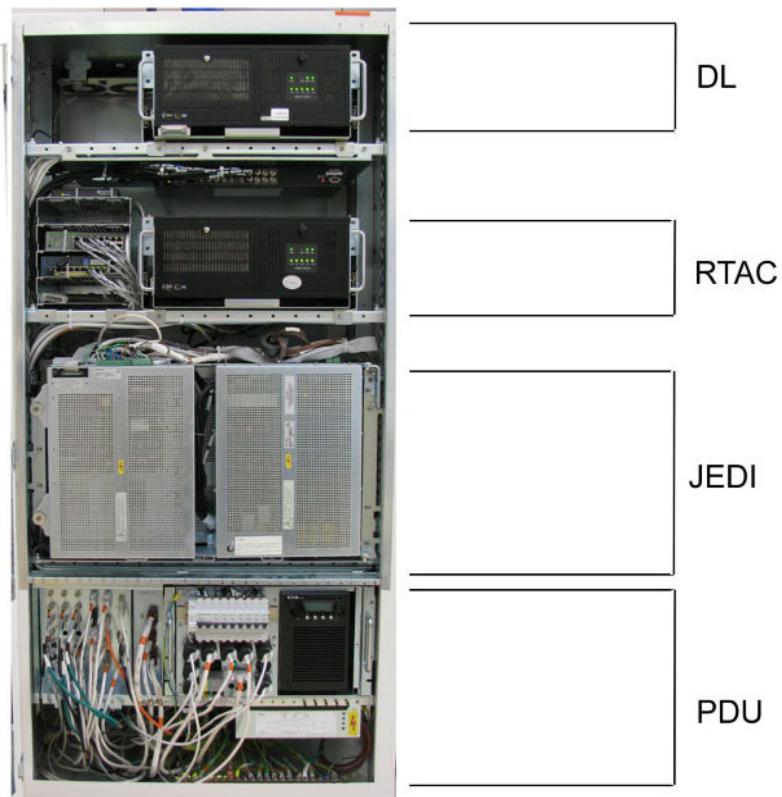


Illustration 2-12: C2 Cabinet

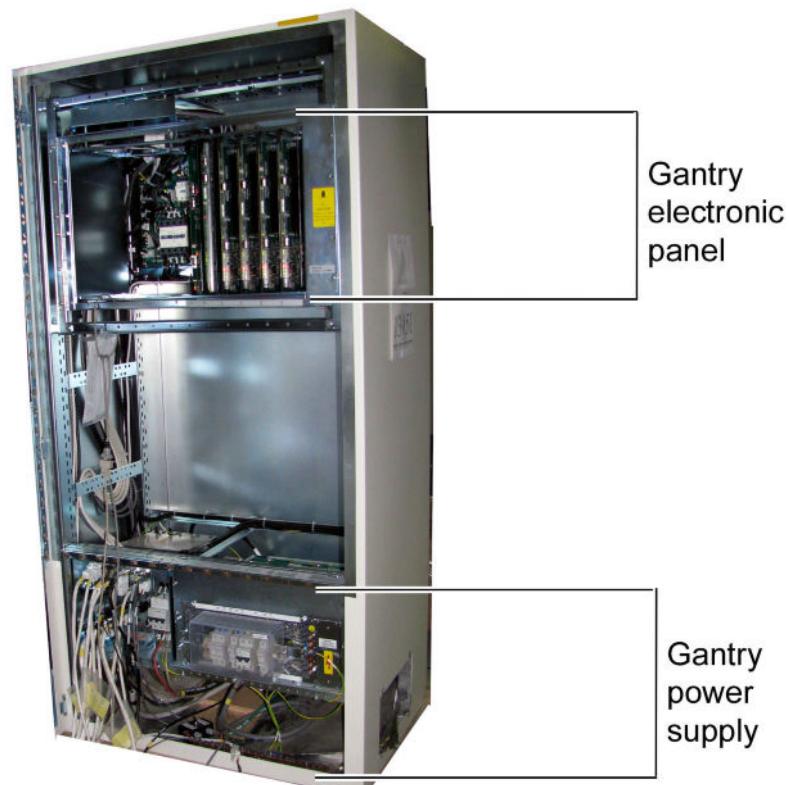


Illustration 2-13: Thermo-con Digital detector chiller



1.2.8 Description of the image monitors suspension

GE provides as option several types of suspensions; alternatively, the customer can install the suspension of his choice ("open suspension"), provided all requirements in the paragraph [Section 1.2.8.3](#) are met.



NOTICE

In the OR configuration of the system, it is mandatory to use a suspension that is compatible with the OR environmental constraints. The suspension provided by GE are not compatible with the OR environment and shall not be used.

1.2.8.1 LCD monitors suspension (without LDM)

The system can be equipped with a suspension for 4 LCD monitors or 6 LCD monitors.

The common type of this suspension is an XT inboard monitor bridge. A monitor frame support receiving 4 or 6 LCD monitors (fixed monitor suspension).

These suspensions are delivered and installed by GE.

1.2.8.2 With the LDM option

For the systems with the LDM option, a specific suspension can be provided, the two backup monitors are mounted on the back of this suspension for faster access in case of failure. For the second LDM, a wall mounting kit can be provided.

1.2.8.3 Open suspension

For Innova IGS systems product with Open Suspension Option, the overhead monitor suspension shall be installed by strictly following the GEHC installation instructions. The manufacturer specifically disclaims any and all liability arising out of or relating to the use or performance of the monitor suspension (including cables), including, without limitation, any liability or claims relating to patient injury, death, or the reliability of such monitors suspension(s).

Where a stand-alone monitor suspension(s) is supplied by the Purchaser of the Innova System, the stand-alone monitors suspension(s) shall comply with the applicable Regulation enforced in the country (eg., when installed in an European Community country, the associated monitors suspension(s) shall be CE marked).

The association of Innova product delivered with Open Suspension Option and the purchaser's (customer) monitors suspension(s), is not covered by Innova product certification.



CAUTION

The Innova System delivered with the Open Monitor Suspension option cannot presume on the mechanical constraints of non-GE monitor(s) suspension(s) introduced in the system.

For further details on Open monitor suspension Pre-installation, please refer to Regulatory Requirements and Pre-Installation Instructions contained in Open Monitor Suspensions - Service Instruction for Installation,5444586-x-1EN contained in the Service manual.

1.2.9 Description of the DL and associated devices

An Innova System uses the DL Digital Imaging system.

- The C1 Cabinet contains:
 1. DL Computer
 2. Firewall
 3. Ethernet switch

- 4. KVM-local
- 5. Video Splitter
- The user area is made of:
 - 1. I/R Receiver / Emitter
 - 2. DL color LCD monitor (19")
 - 3. Keyboard,
 - 4. Mouse
 - 5. Keypads for user dialogue
 - 6. Video station:
 - Hi-Brite monitor in a user area
 - VCR in option

Illustration 2-14: Innova User Area devices



Remote control

Keypad

Innova VCIM with DL keyboard console



DL color LCD monitor



Post processing color LCD monitors
(Options)



Repeater B&W LCD monitors (Options)



Advantage
windows
monitor



Centricity
CA1000
monitor



Large Display Monitor (Option)

1.2.10 AW workstation

AW workstation option is composed of a workstation, 1 or 2 monitors 19" flat panel in the Control Room and 1 monitor (flat panel) fixed on suspension (option) and 1 video switcher. Check Innova Installation Service Manual for switcher installation.

1.2.11 CENTRICITY CA1000 option

Refer to :*Centricity Cardiology CA 1000 V2.0 Preinstallation Guide* in the OEMs of the Innova IGS 520, Innova IGS 530, Innova IGS 540 service manual.

1.2.12 Injectors

NOTE: The following inventory lists all injectors. For the list of injectors compatible with your system configuration, refer to the Operator Manual.

(For Omega Table) the recommended injectors are:

- LF Angiomat Illumina (pedestal version) or
- MedRad Mark V ProVis (table/rack mount version) or
- MedRad Avanta (pedestal version) or
- MedRad Avanta (table mount version) or
- MedRad Mark 7 (table mount version without extension cable) or
- MedRad Mark 7 (table mount version with extension cable) or
- MedRad Mark 7 (pedestal mount version without extension cable) or
- Acist CVI (pedestal version) or
- Acist E2000 Voyager (pedestal version) or
- Acist CVI (table mount version).

(For Tilting Table) the recommended injectors are:

- MedRad Mark V ProVis (table/rack mount version) or
- MedRad Avanta (table/rack mount with rack connected to table foot or pedestal version) or
- MedRad Mark 7 (pedestal mount version without extension cable) or
- Acist CVI (pedestal version)

NOTE: With Tilting Table:

- ACIST CVI Table mount injector is not authorized.
- Table Mounted Injectors are not allowed to mount on the OR Table as they are not compatible:
 - MEDRAD Mark V Provis
 - MEDRAD Avanta

1.2.13 Fluoro UPS

The Innova system can be protected with an optional Fluoro UPS.



FLUORO UPS IS MANDATORY FOR OR TABLE.

There are two types of Fluoro UPS:

- UL for North America and other 480V – 60Hz countries.
- CE for Asia and Europe

Illustration 2-15: Fluoro UPS



UPS UL

NOTE: Refer to the UPS vendor Service Operating Manual for more details:

- [UL Model: UPS Option - Operating Manual SG-UL Series - 20 kVA](#)
- [CE Model: UPS Option - Operating Manual SitePro 20 kVA \(series 8\)](#)

Illustration 2-16: UPS IF box



UPS IF BOX

1.2.14 Volcano IVUS s5i Option

NOTE: For IVUS Rev 3 components, refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

1.2.14.1 Purpose

Volcano is a manufacturer of Intravascular Ultrasound Systems (IVUS), which are used in the cath lab. GE has a strategic agreement with Volcano. GE will be selling and servicing some of the Volcano products.



FOR OR TABLE, IVUS ACCESSORIES MUST BE INSTALLED ON A CART

1.2.14.2 Product Description

The Volcano S5I GE systems are dedicated equipments for IntraVascular UltraSound imaging (S5I GE) and Fractional Flow Reserve (FFR) evaluation, designed as add on equipment to Vascular Imaging systems. The S5I GE equipment is also capable of receiving patient data from the Innova system.

The Volcano S5I GE Hardware configuration consists of:

- A computer (S5I GE CPU) connected to a monitor (S5I GE Monitor), both located in the Control Room,

- Patient Interface Modules as the S5I GE PIM, the PIMr for rotational catheters and Pimette for the FFR catheter, located in the Procedures Room,
- Various user interface units to control the system
 - S5I GE Control Station,
 - S5I GE Joystick,
 - S5I GE Touchpad Controller
 - S5I GE Keyboard & mouse pad installed in the Control Room
- Optional Image Printer, located in the Control Room is available for the S5I GE system

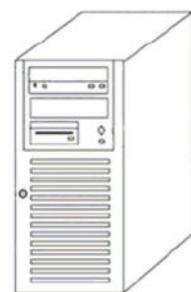
The IVUS integration to Innova, involves further components :

- Innova Central (Touchscreen) as optional control station,
- Video Signal Switch to route Workstation and S5I GE video signals,
- Procedures Room monitor shared by Workstation (AW or CA1000) and S5I GE operations,
- Network cable (S5I GE to Innova system),
- Grounding cable (S5I GE Isolation Transformer to C1 Cabinet ground bar),

Illustration 2-17: IVUS Rev 2 components



IVUS Safety Isolation Transformer



IVUS CPU



IVUS Patient Interface module



IVUS Control Console



IVUS Video Switch



IVUS Joystick



IVUS printer



IVUS Control Room Monitor



IVUS Touchpad Control



IVUS PIMr



IVUS FFR Pimette

Illustration 2-18: IVUS Rev 2+ components



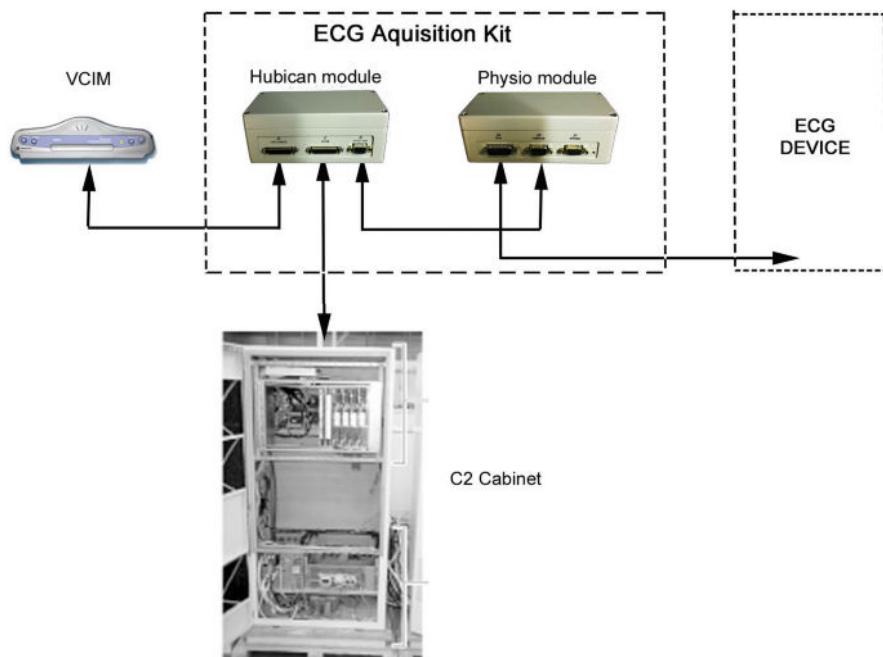
1.2.15 ECG Acquisition kit Option

The ECG Acquisition kit allows for the connection of an ECG device (such as GE ECG devices MacLab, CardioLab or ComboLab) to the Innova system (C2 cabinet). The ECG Acquisition kit hardware consists of a Hubican and Physio modules and their associated cables.

There are two possible configurations for the ECG Acquisition kit:

- ECG devices installed in Control Room (GE ECG devices like MacLab, CardioLab or ComboLab)
- ECG devices installed in Exam Room (Non-GE ECG device)

Illustration 2-19: ECG Acquisition kit layout



1.2.16 Vivid E9 Diagnostic Ultrasound System Option

The VIVID E9 ultrasound unit is a high performance digital ultrasound imaging system with total data management.

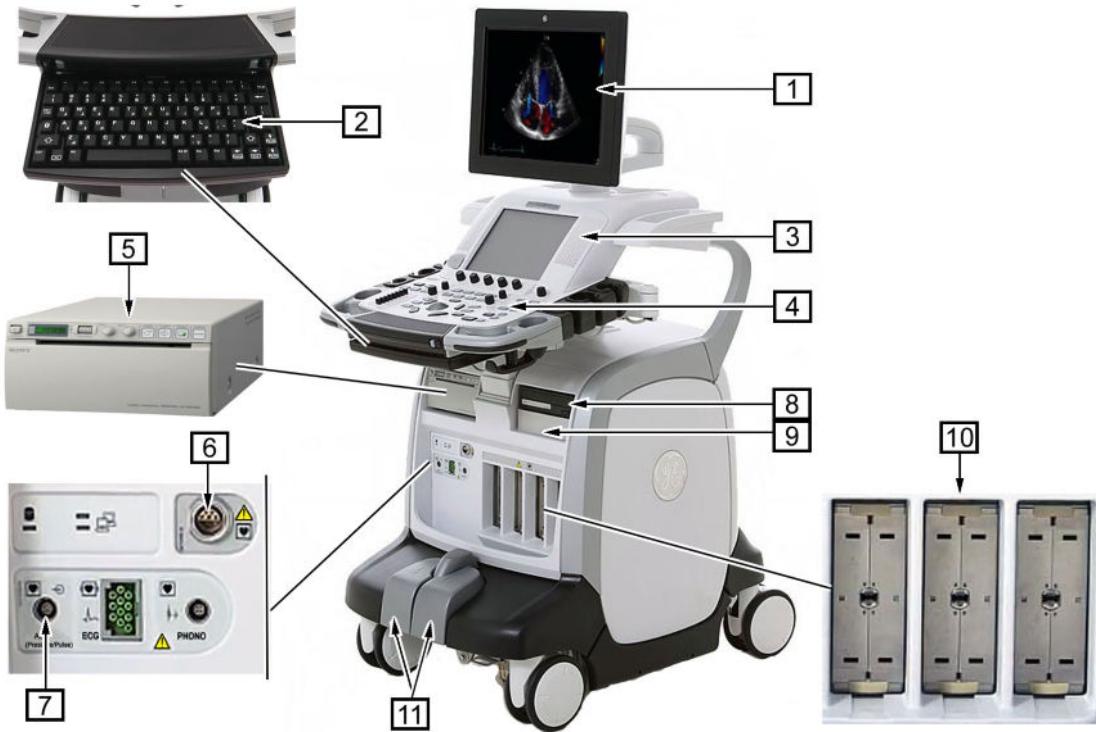
It is installed in the vascular Patient Room's environment as the clinical procedure requires. After the procedure is completed, it might be moved out of the room for other application purposes.

The system provides image generation in 4D, 2D (B) Mode, Color Doppler, Power Doppler (Angio), MMode, Color M-Mode, PW and CW Doppler spectra, Tissue Velocity imaging, Advanced Strain and Contrast applications. The fully digital architecture of the VIVID E9 unit allows optimal usage of all scanning modes and probe types, throughout the full spectrum of operating frequencies.

The Signal runs from the Probe Connector Panel to the Front End, then over to the Back End Processor and finally to the monitor and peripherals.

System configuration is stored on the hard drive in the Back End Processor. All necessary software is loaded from the hard drive on power up.

Illustration 2-20: VIVID E9 major components



VIVID E9 major components, refer to [Illustration 2-20](#):

- 1 Main Monitor
- 2 Alphanumeric Keyboard
- 3 Upper panel with Stereo Loudspeakers and Touch Monitor
- 4 Operator Panel
- 5 B/W Video Printer
- 6 Doppler (PEDOF) Connector
- 7 Patient I/O
- 8 DVD Player
- 9 DVD Recorder
- 10 Probe Connectors
- 11 Brake Pedals

For Product Identification labelling, please refer *VIVID E9 Service Manual GA091568TPH* Section 1-5

For further details please refer *VIVID E9 Service Manual GA091568TPH* Section 1-2-4.

1.2.17 Large Display Subsystem option

The Innova system can integrate a Large Display solution to:

- See images larger at full IQ with greater flexibility in monitor distance in the procedure room
- Display multiple video images simultaneously at different sizes based on stage of workflow
- Conveniently switch operator defined video layouts at different points in procedure workflow

The Large Display Subsystem option is addressing: 100, 110, 220, 230, 240 VAC (phase, neutral and ground) +/- 10% 50Hz +/-3Hz and 60Hz +/-3Hz countries

Large Display Monitor Configuration:

- Large Display Monitor: 58" Eizo color
- Black and white monitors: 19" Eizo monitor

1.3 Dimension Drawings

Refer to this section for the dimensional drawings of the components of the Gantry and Patient Tables sub-systems. These components include:

Gantry, Omega Tables, Tilting Tables, C2 Cabinet, C1 Cabinet, and Chillers. In addition, refer to this section for Gantry/Table relative position drawings.

Table 2-1:

TITLE	ILLUSTRATION
Gantry Dimensions: - Side View	Illustration 2-21
Gantry Dimensions: - Top View	Illustration 2-22
Gantry Dimensions: - Front View	Illustration 2-23
Omega IV Compact Patient Table Dimensions	Illustration 2-24
Omega IV Table Interference Regions (Innova IGS 520)	Illustration 2-25
Omega V Long Patient Table Dimensions	Illustration 2-26
Omega V Long Patient Table Interference Regions	Illustration 2-27
Tilting table Dimensions	Illustration 2-28
Tilting Table Interference Regions	Illustration 2-29
Patient Table side clearance (CPR access)	Illustration 2-30
Table Head Extender	Illustration 2-31
Gantry and Omega IV Compact Patient Table Relative Positions (Innova IGS 520)	Illustration 2-32
Gantry and Omega V Long Patient Relative Positions	Illustration 2-33
C2 Cabinet Dimensions	Illustration 2-34
C1 Cabinet Dimensions	Illustration 2-35
Dimension Diagram for X-Ray Tube Chiller 4100	Illustration 2-36
Floor Space Diagram for X-Ray Tube Chiller 4100	Illustration 2-37
Detector Chiller Thermo-Con Dimensions & Orientation	Illustration 2-38
Fluoro UPS UL Layout (Optional)	Illustration 2-39
Fluoro UPS CE Layout (Optional)	Illustration 2-40
UPS IF Box (Optional)	Illustration 2-41
3 kVA UPS	Illustration 2-42
EMI Filter Box (CE Only)	Illustration 2-43
Gas box outlets Omega IV	Illustration 2-44
Gas box outlets Omega V	Illustration 2-45
DL Keypad Dimensions	Illustration 2-46
DL Image Monitor Dimensions	Illustration 2-47
VCR Mounting Holes Location	Illustration 2-48
ECG Acquisition Device Modules	Illustration 2-49
Large Display cabinet dimensions (Optional)	Illustration 2-50
Large Display suspension dimensions (Optional)	Illustration 2-51

Illustration 2-21: Gantry Dimensions: - Side View

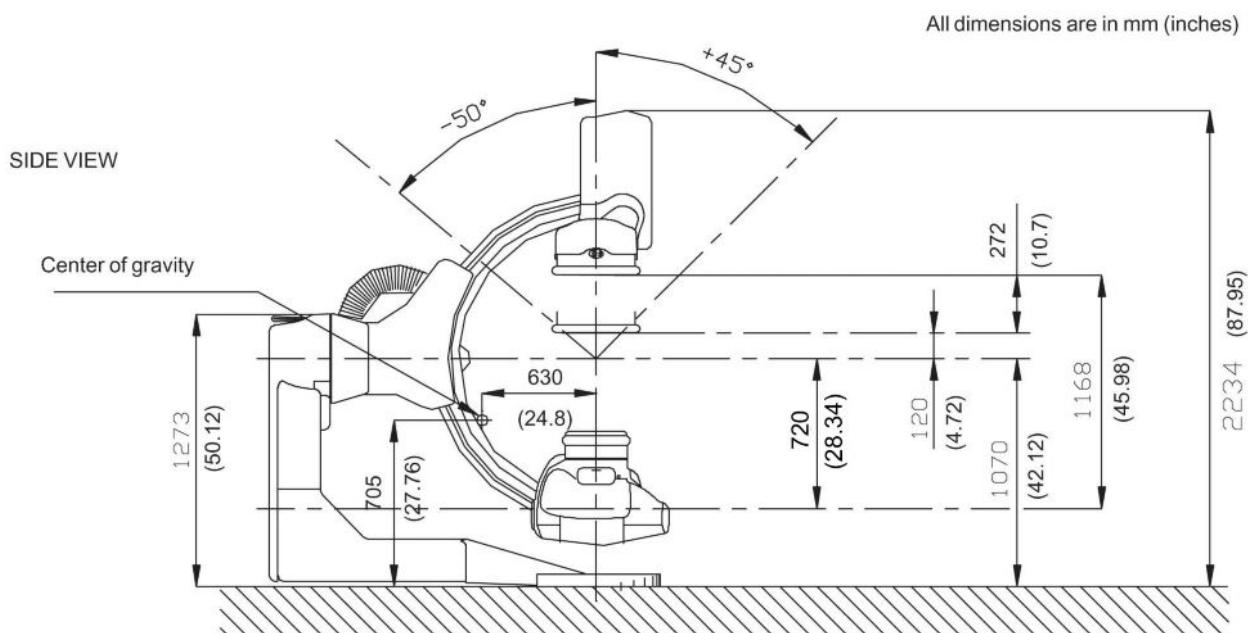


Illustration 2-22: Gantry Dimensions: - Top view

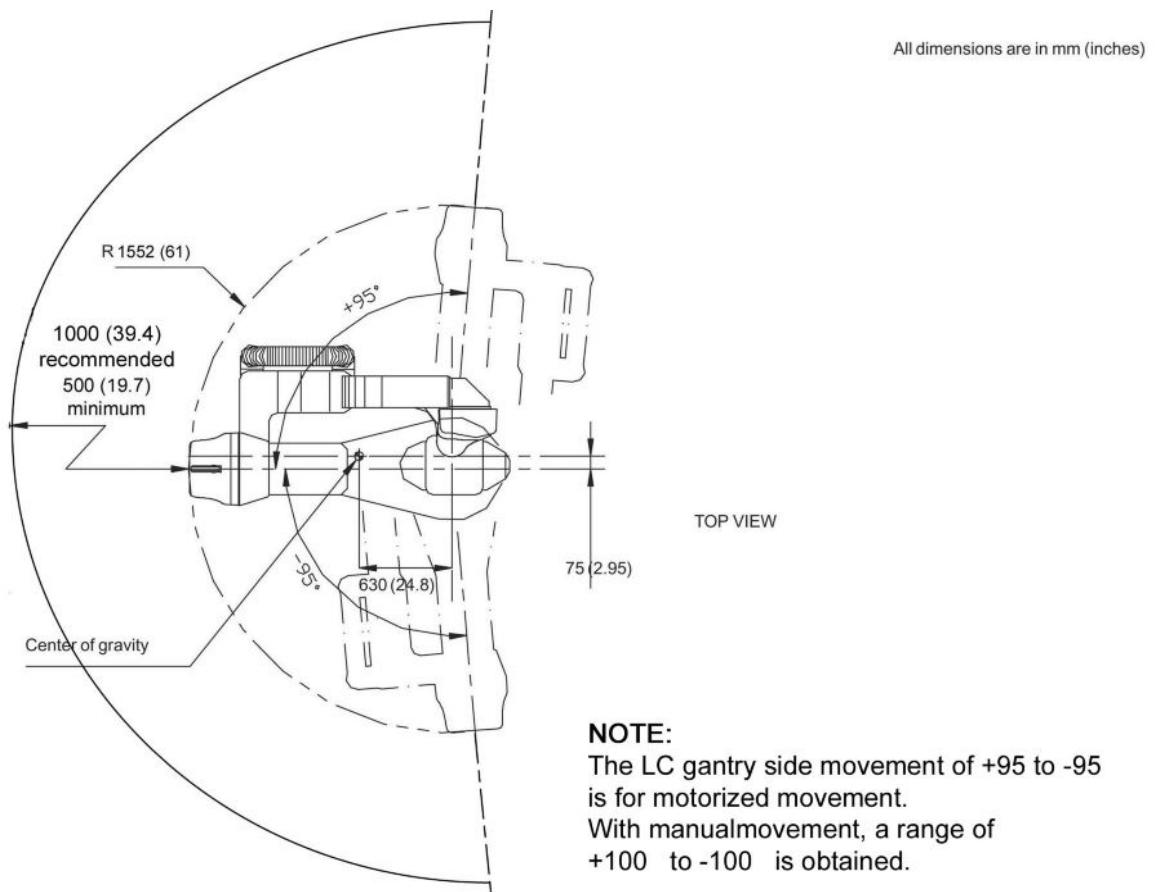


Illustration 2-23: Gantry Dimensions: - Front view

All dimensions are in mm (inches)

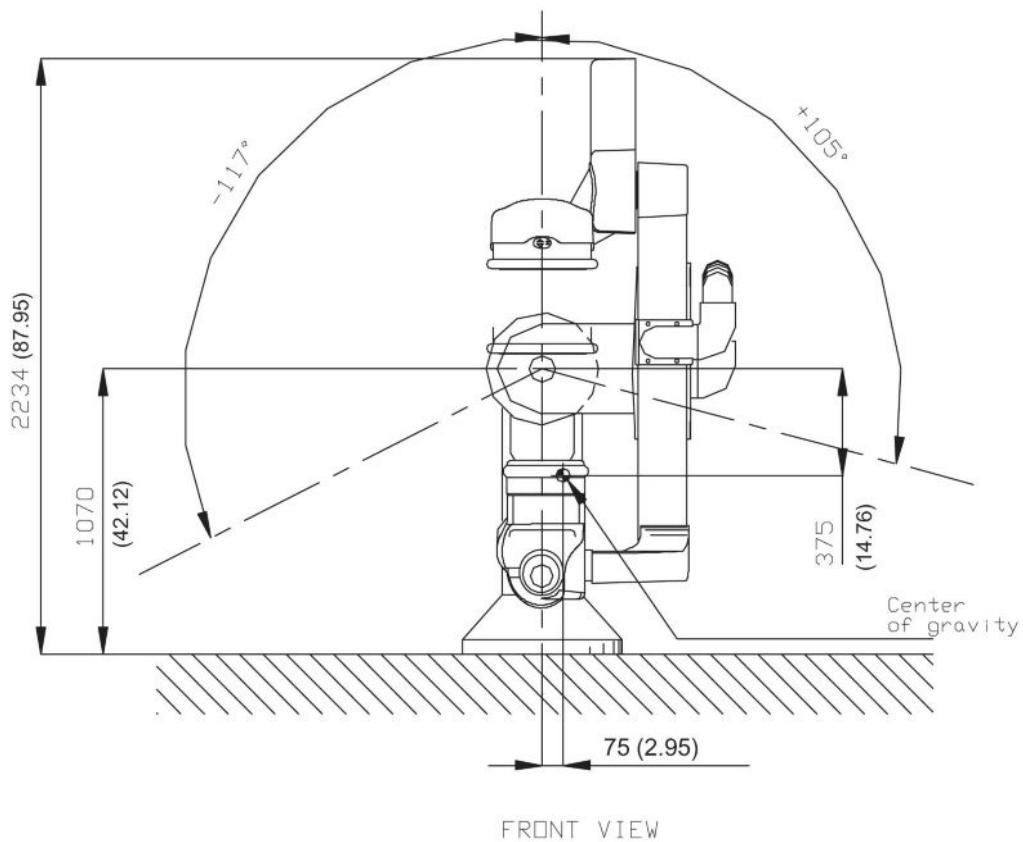


Illustration 2-24: Omega IV Compact Patient Table Dimensions

All dimensions are in mm (inches)

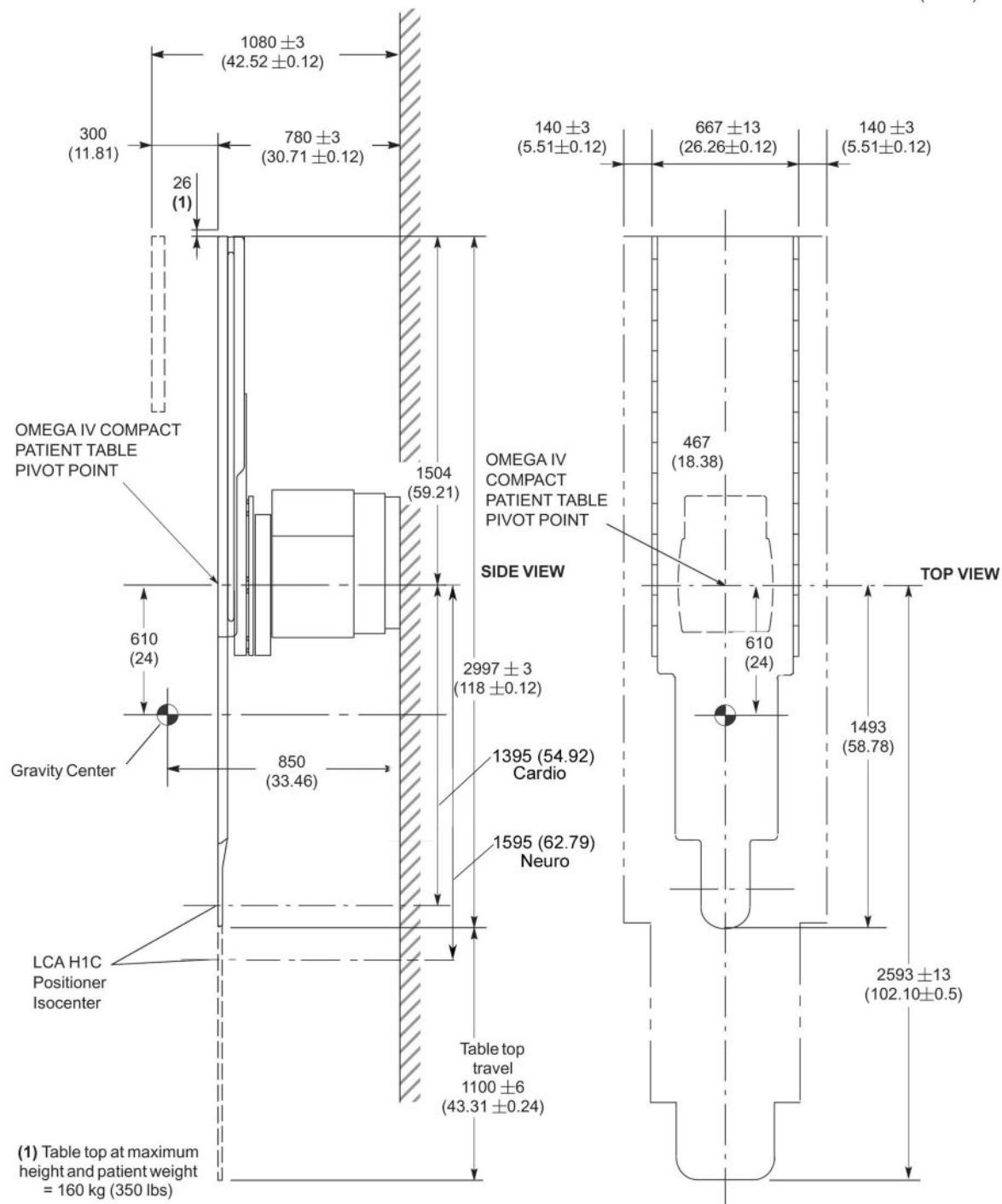


Illustration 2-25: Omega IV Compact Patient Table Interference Regions (Innova IGS 520)

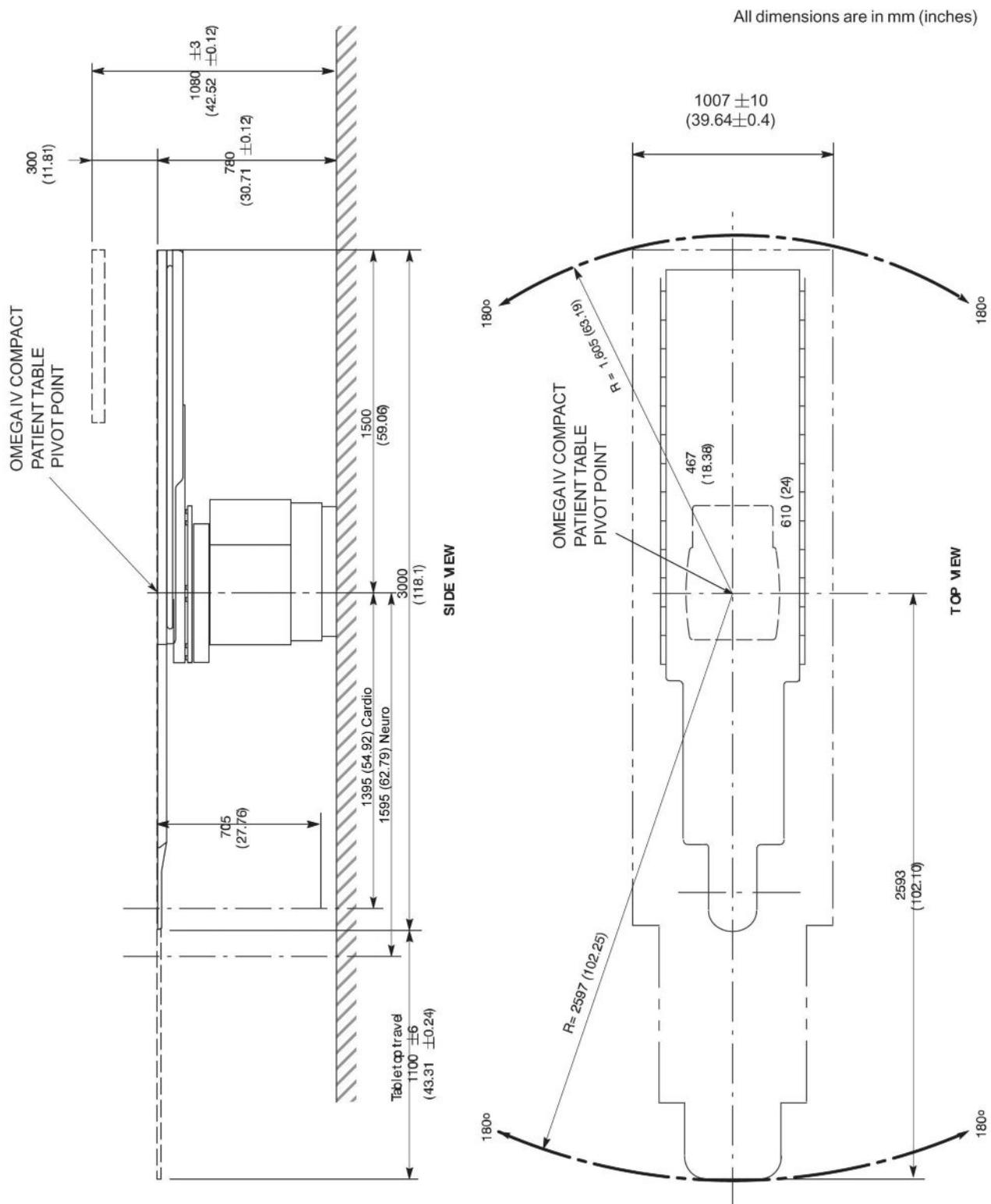
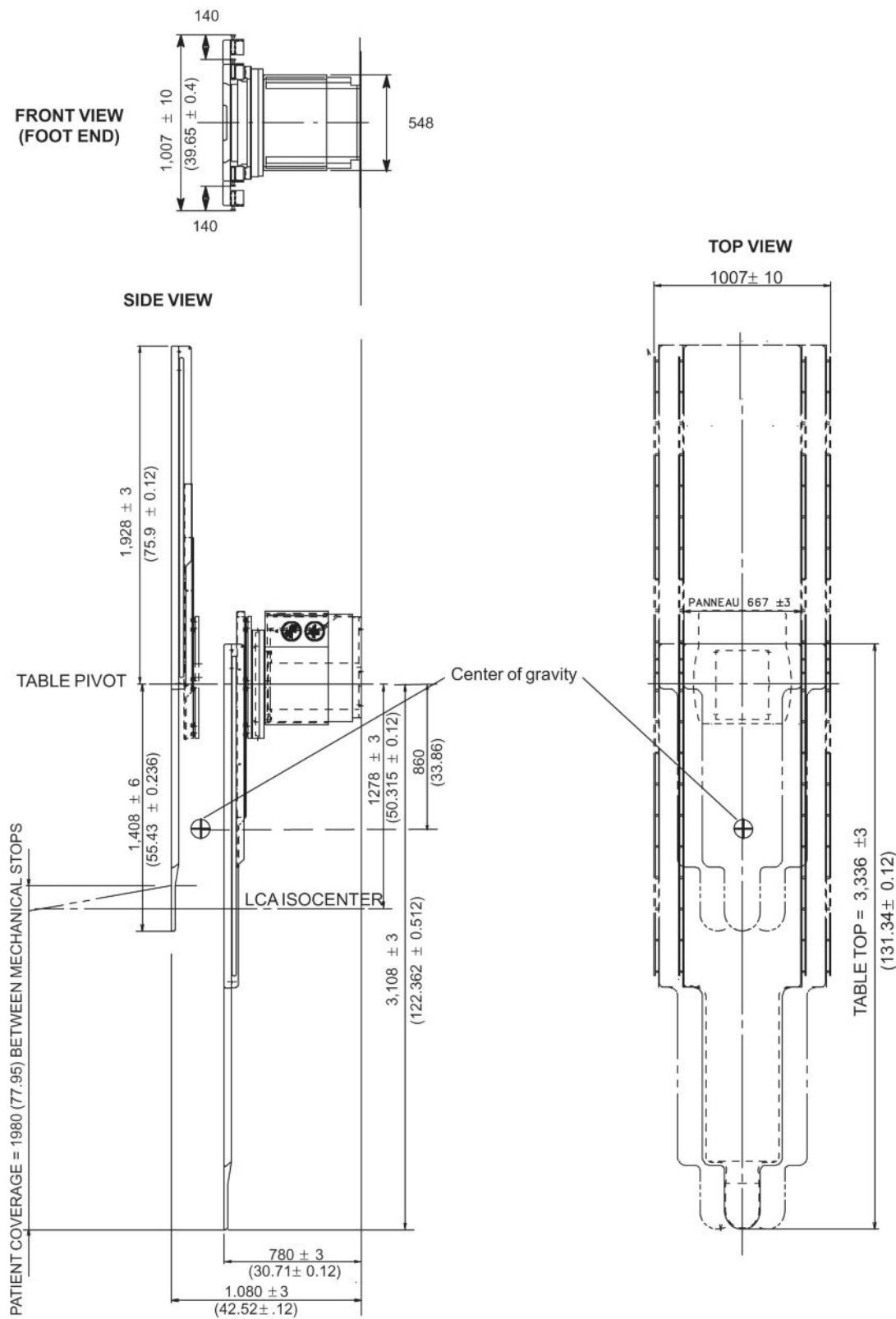


Illustration 2-26: Omega V Long Patient Table Dimensions



All dimensions are in mm (inches)

Illustration 2-27: Omega V Long Patient Table Interference Regions

All dimensions are in mm (inches)

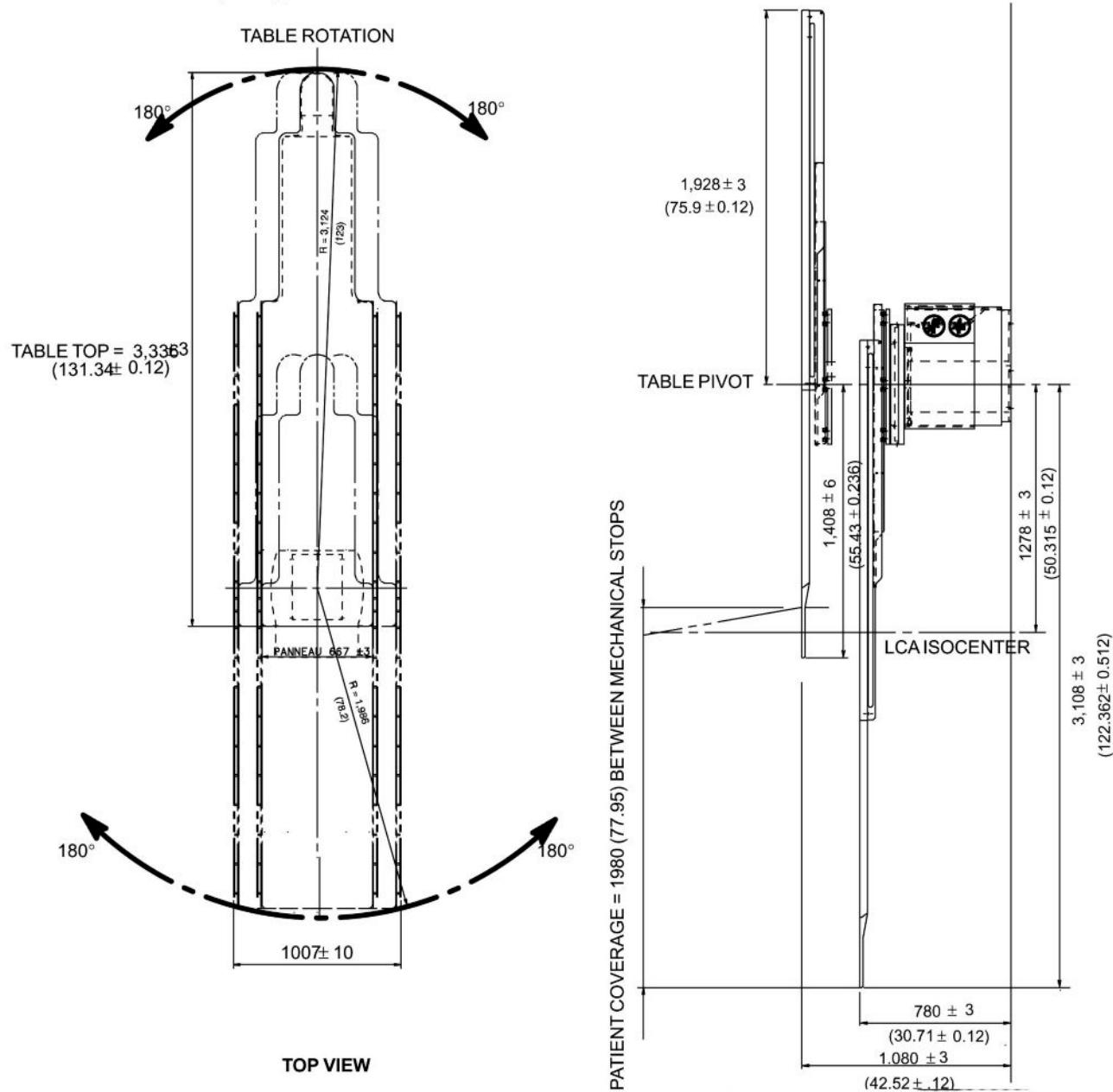
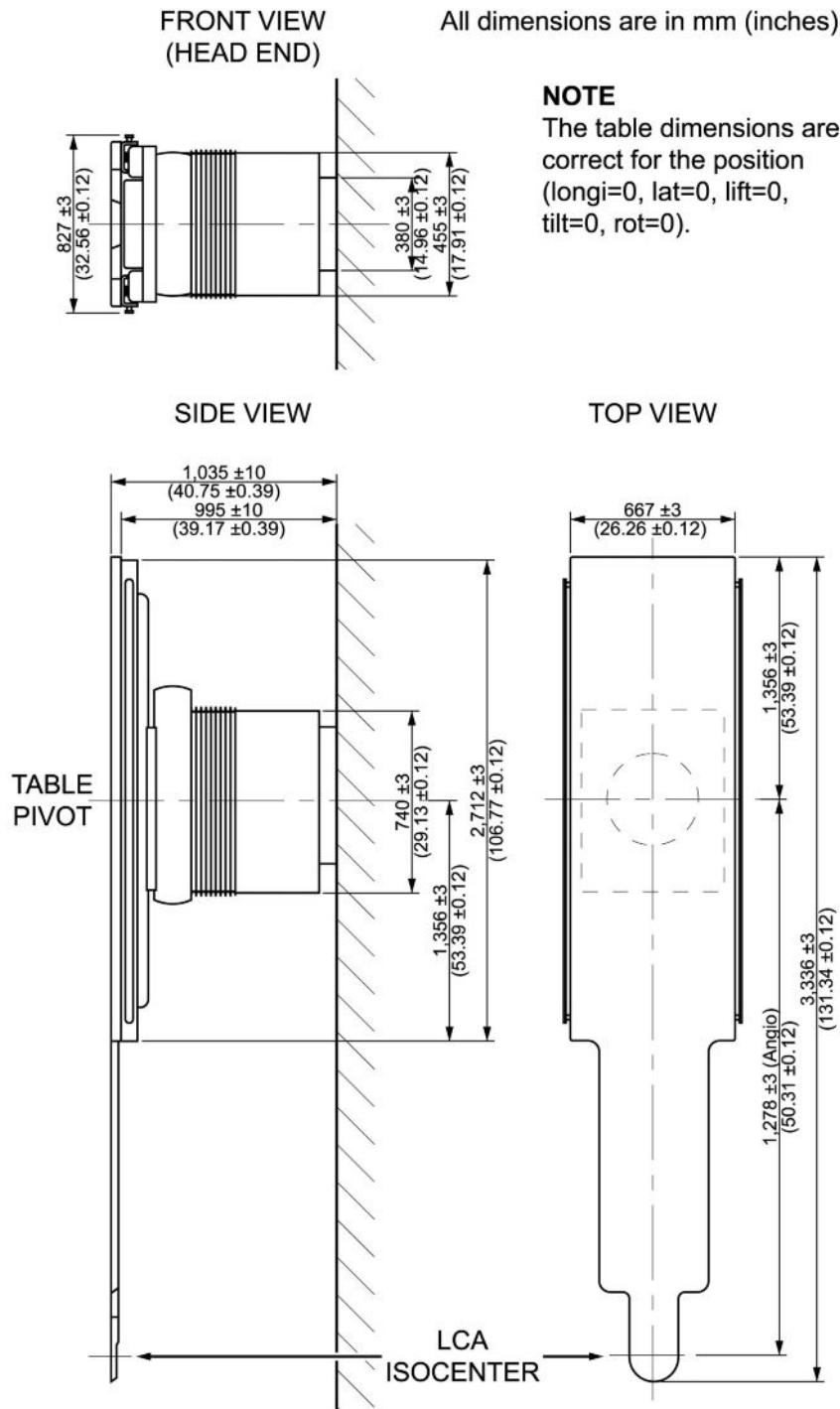


Illustration 2-28: Tilting table Dimensions



NOTE: With a Tilting table, the minimum distance from table pivot to the medical Gas Box is 600 mm and the maximum dimensions of the medical Gas Box are :

- height = 305 mm
- width = 250 mm
- length = 500 mm

Illustration 2-29: Tilting table Interference Regions

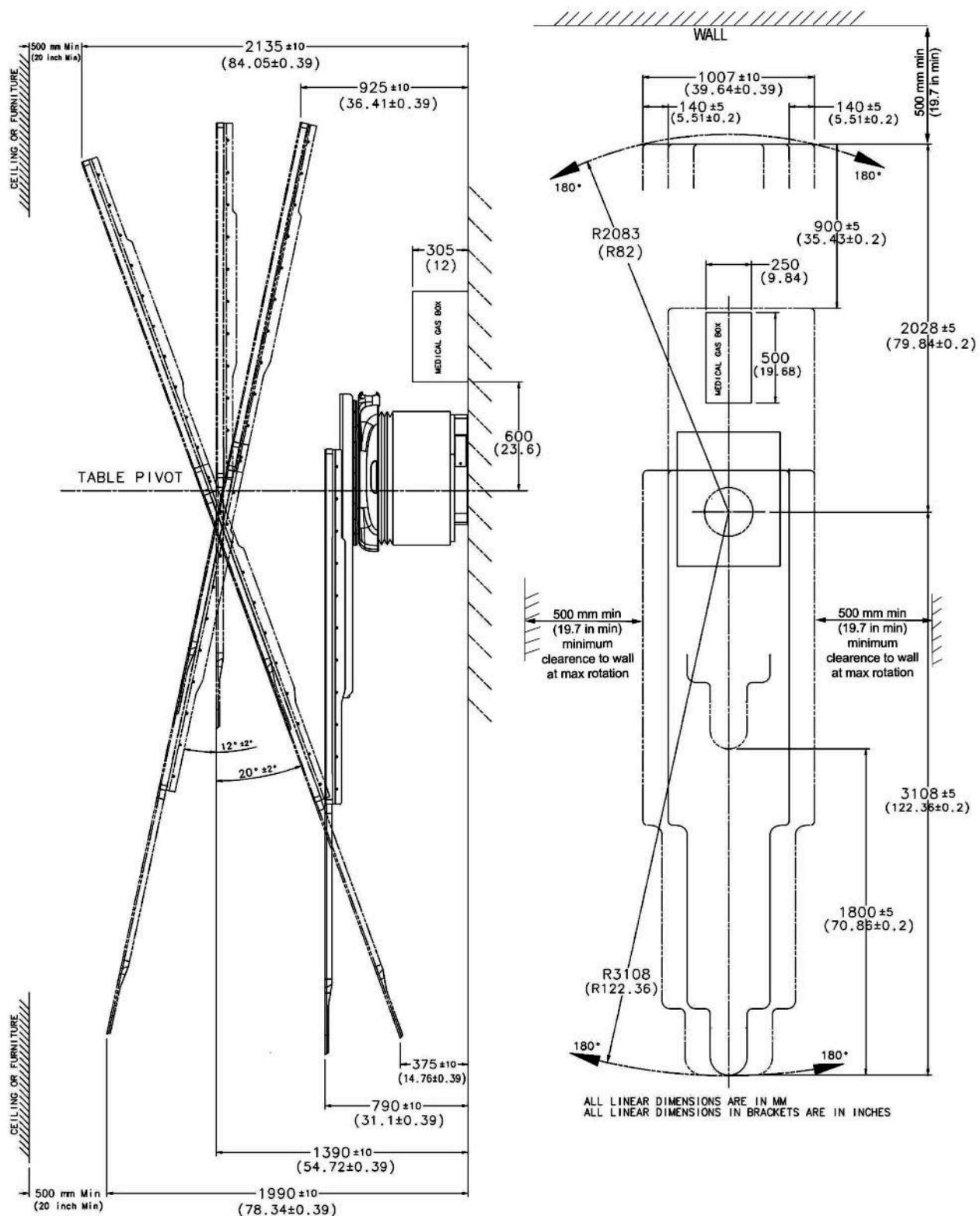


Illustration 2-30: Patient Table side clearance (CPR access)

All dimensions are in mm (inches)

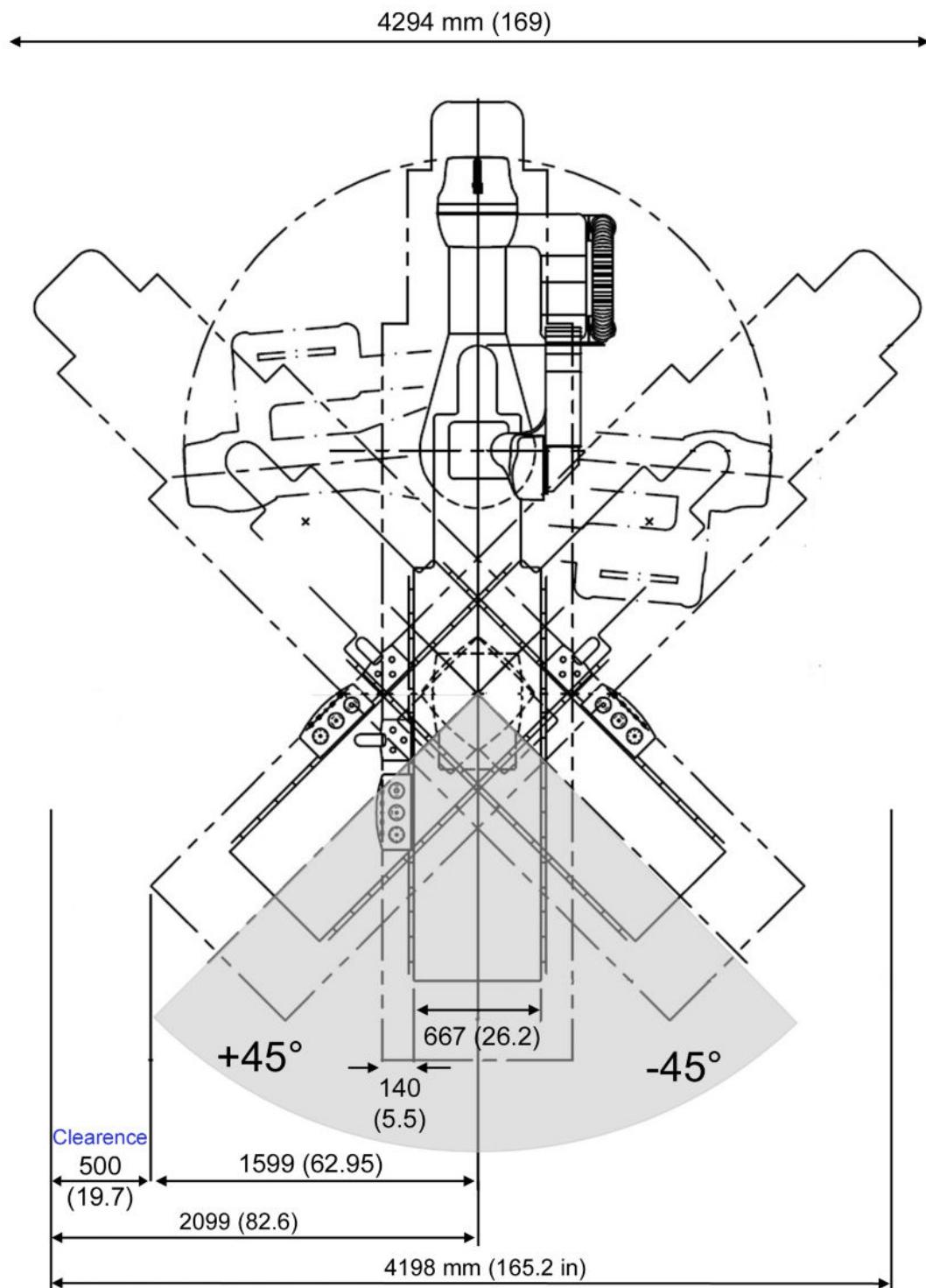


Illustration 2-31: Table Head Extender

All measurements are in mm (inches)
Based on drawing 5262690ADW

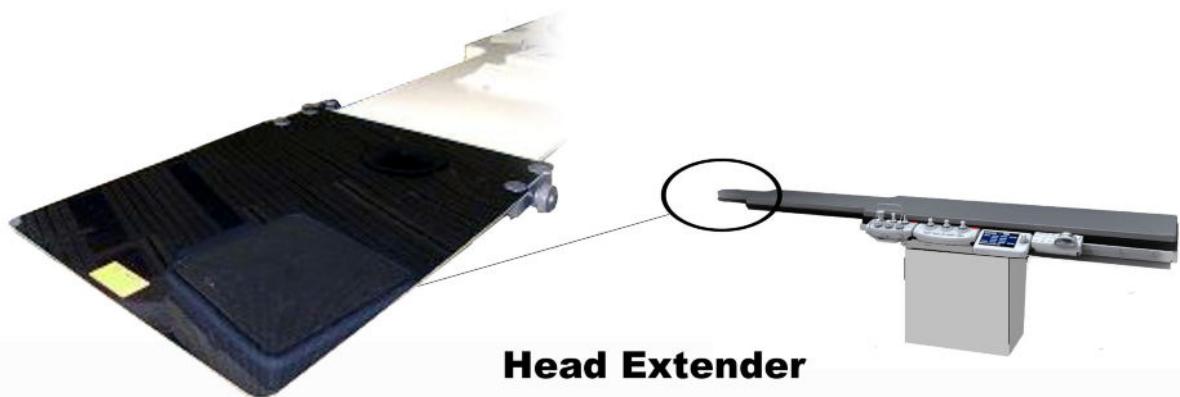
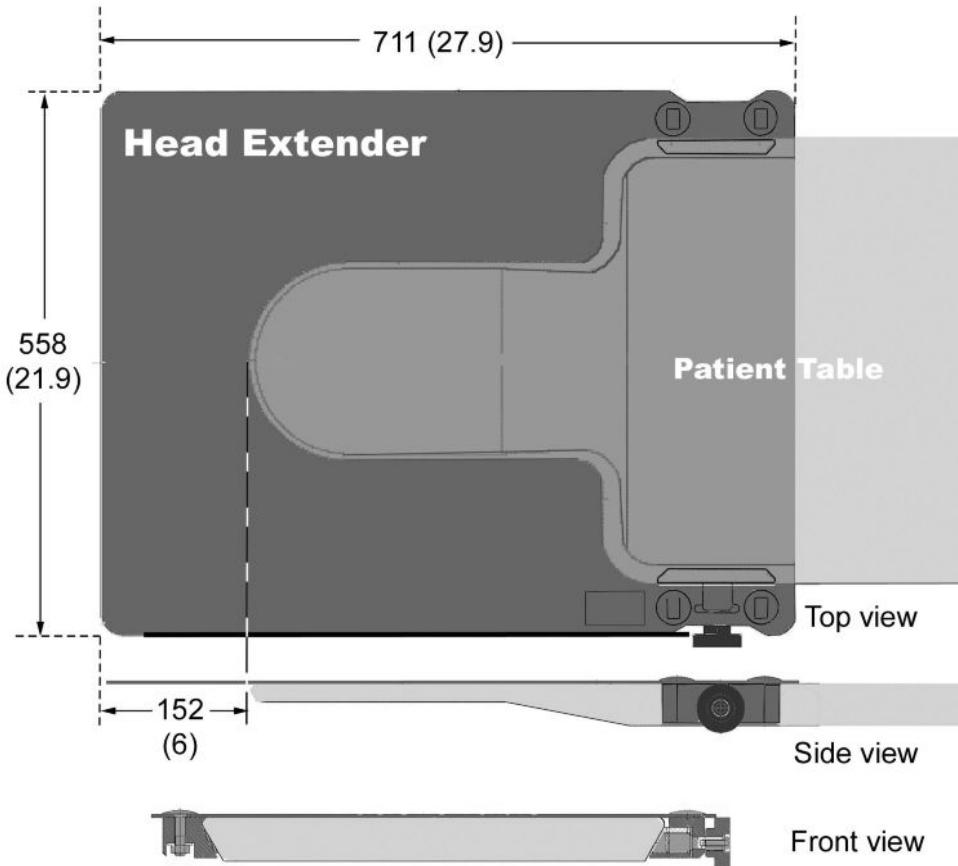
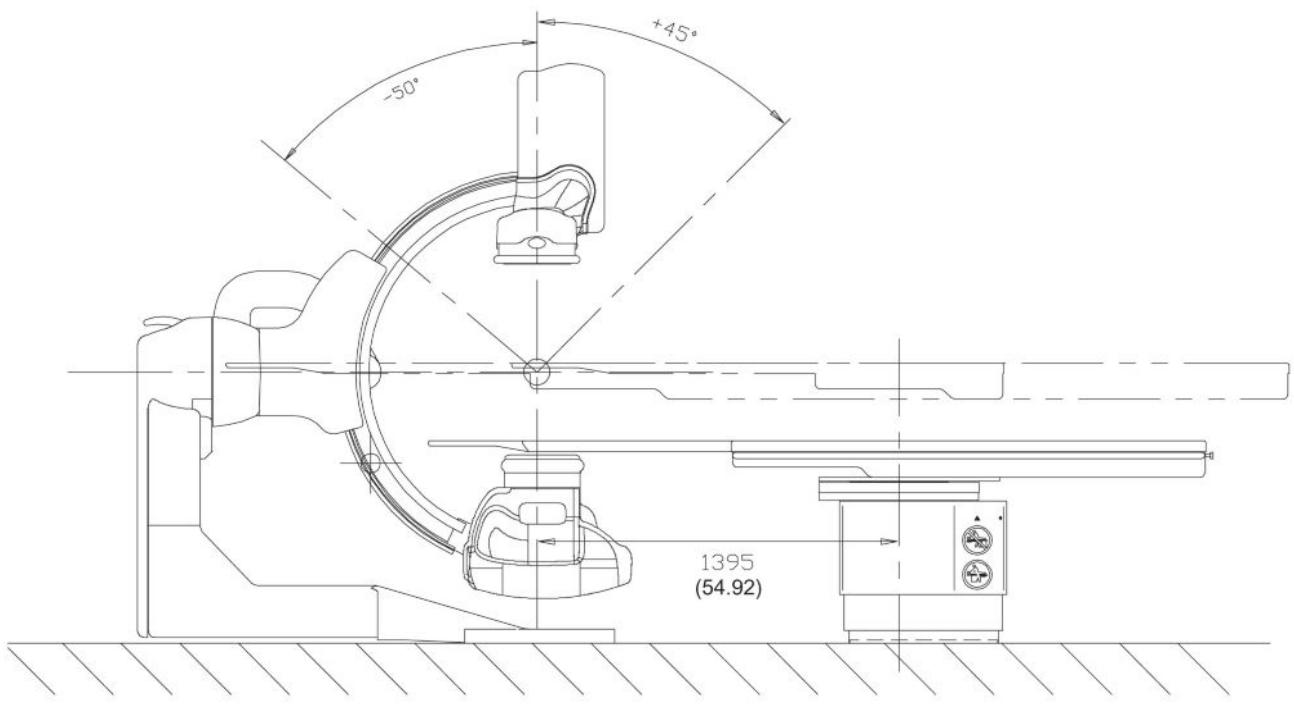


Illustration 2-32: Gantry and Omega IV Compact Patient Table Relative Positions (Innova IGS 520) - side view

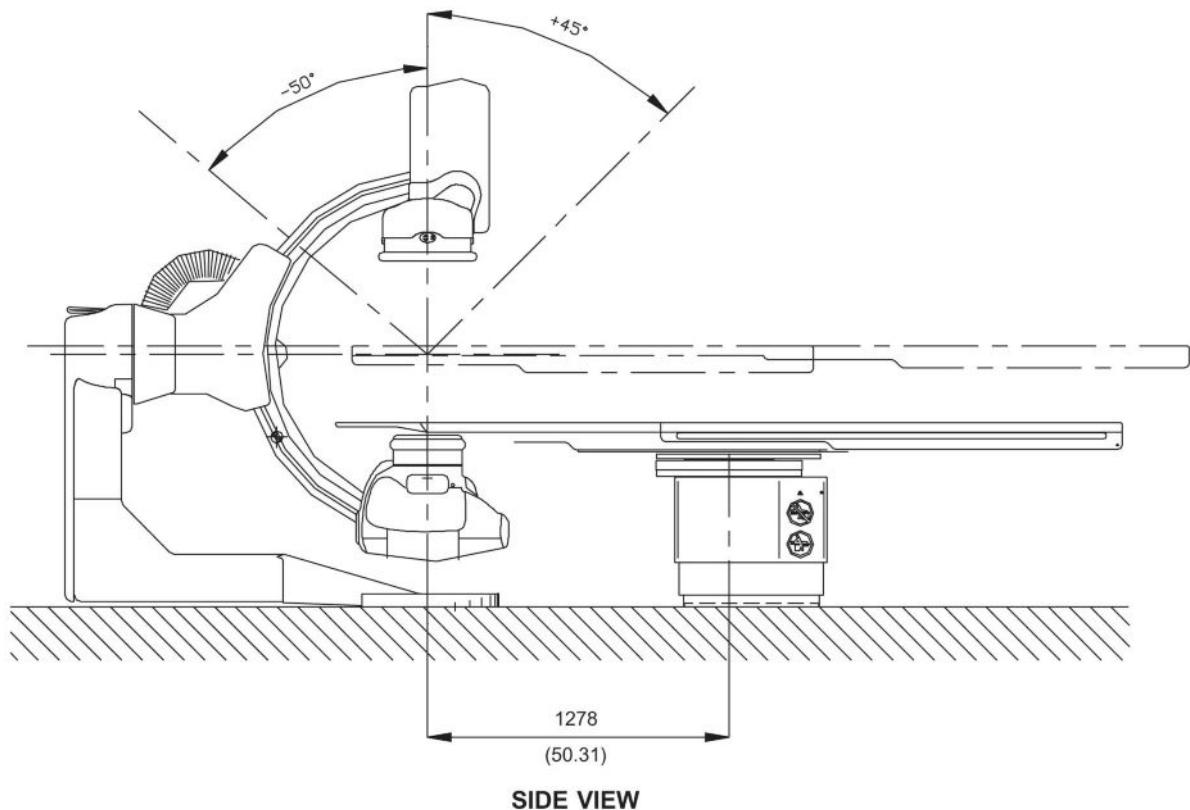
All dimensions are in mm (inches)



SIDE VIEW

Illustration 2-33: Gantry and Omega V Long Patient Table Relative Positions - side view

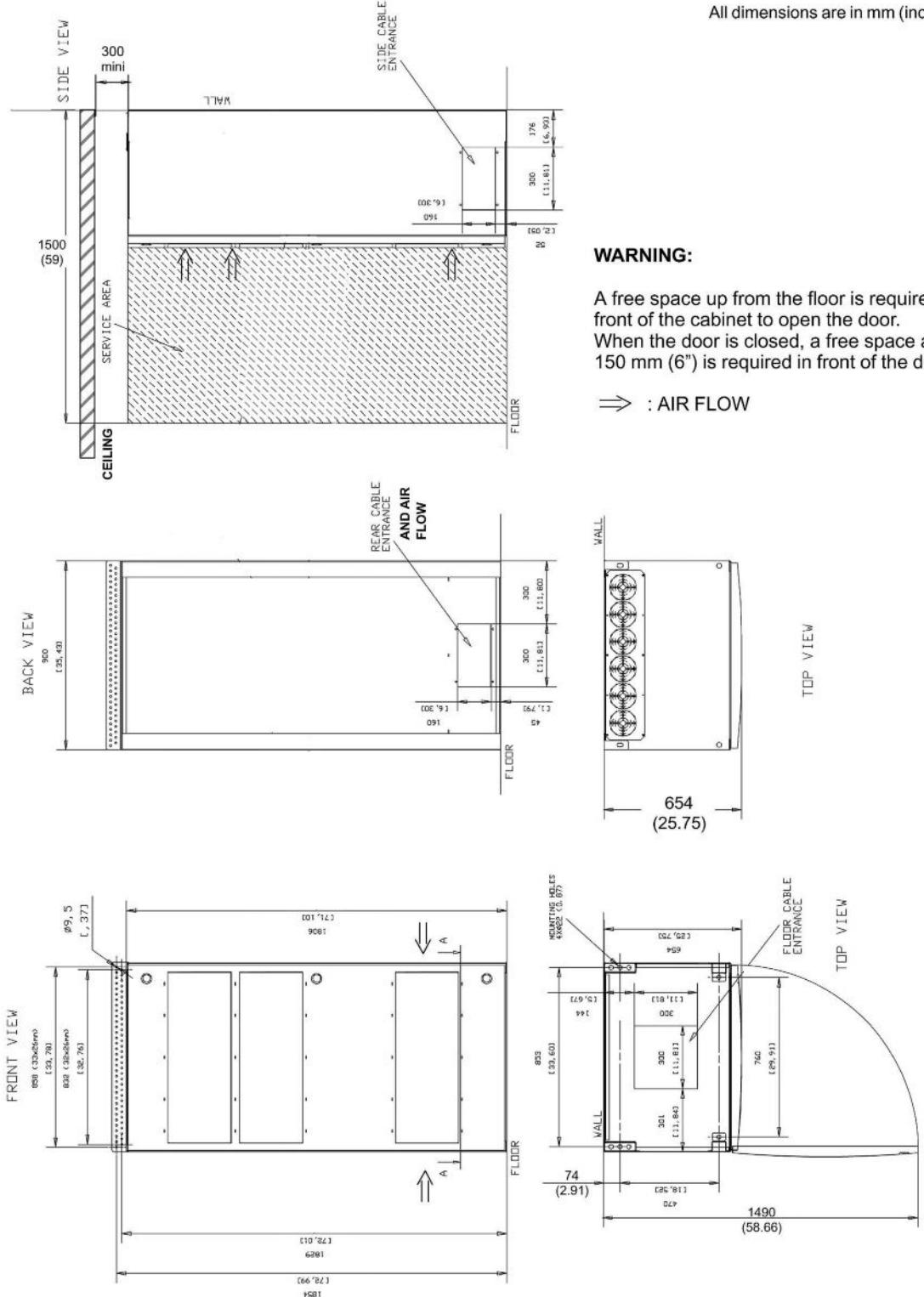
All dimensions are in mm (inches)



SIDE VIEW

Illustration 2-34: C2 Cabinet Dimensions

All dimensions are in mm (inches)

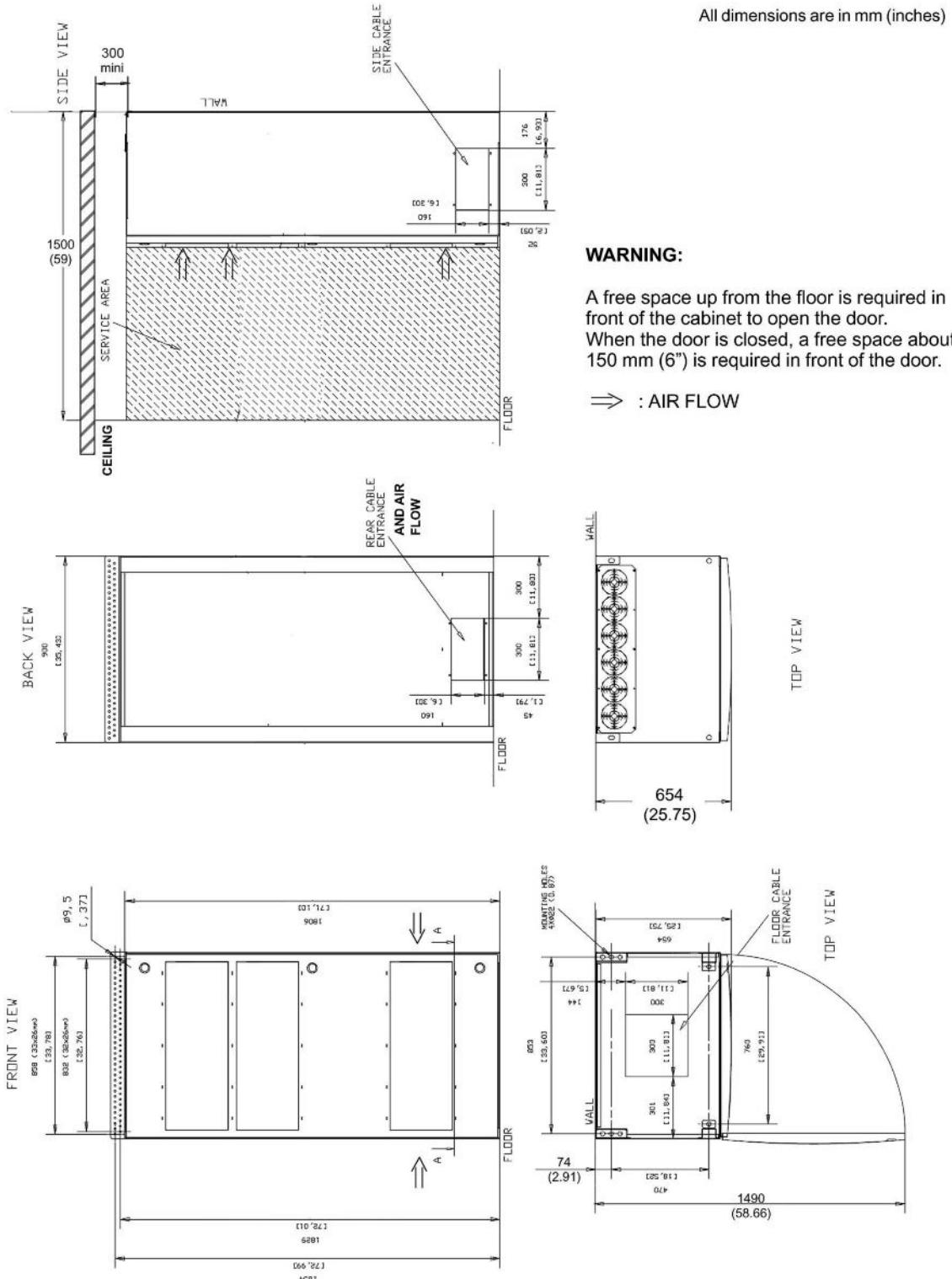


WARNING:

A free space up from the floor is required in front of the cabinet to open the door.
When the door is closed, a free space about 150 mm (6") is required in front of the door.

⇒ : AIR FLOW

Illustration 2-35: C1 Cabinet Dimensions

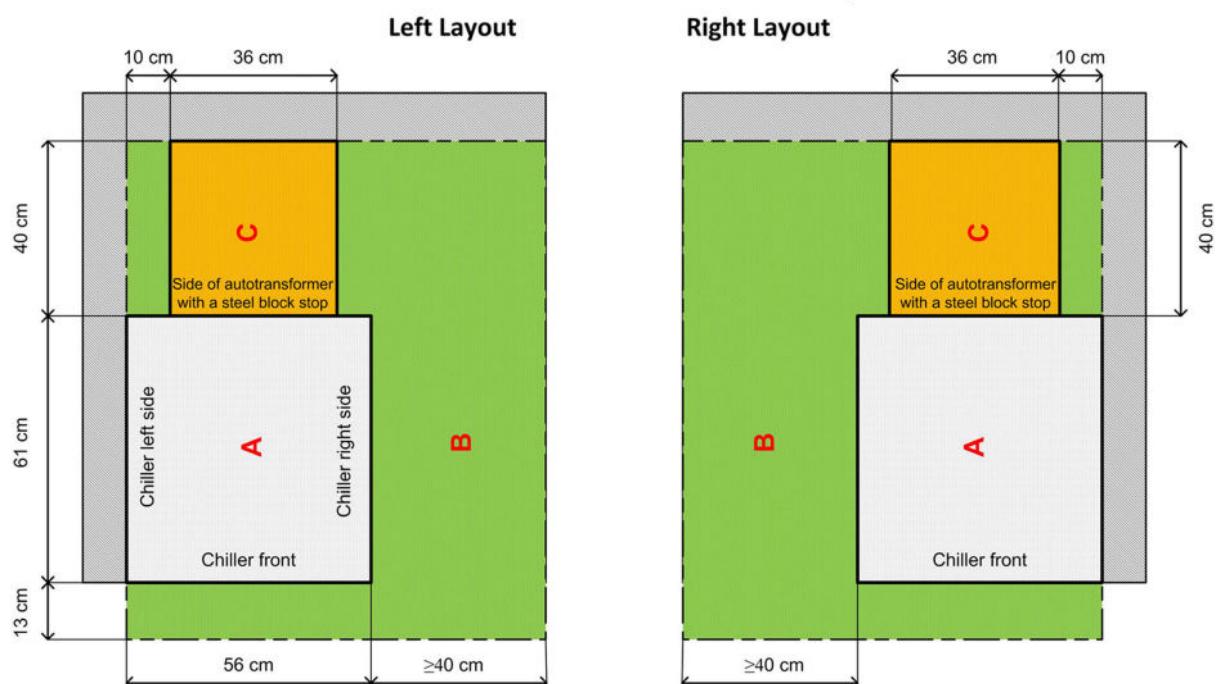


WARNING:

A free space up from the floor is required in front of the cabinet to open the door.
When the door is closed, a free space about 150 mm (6") is required in front of the door.

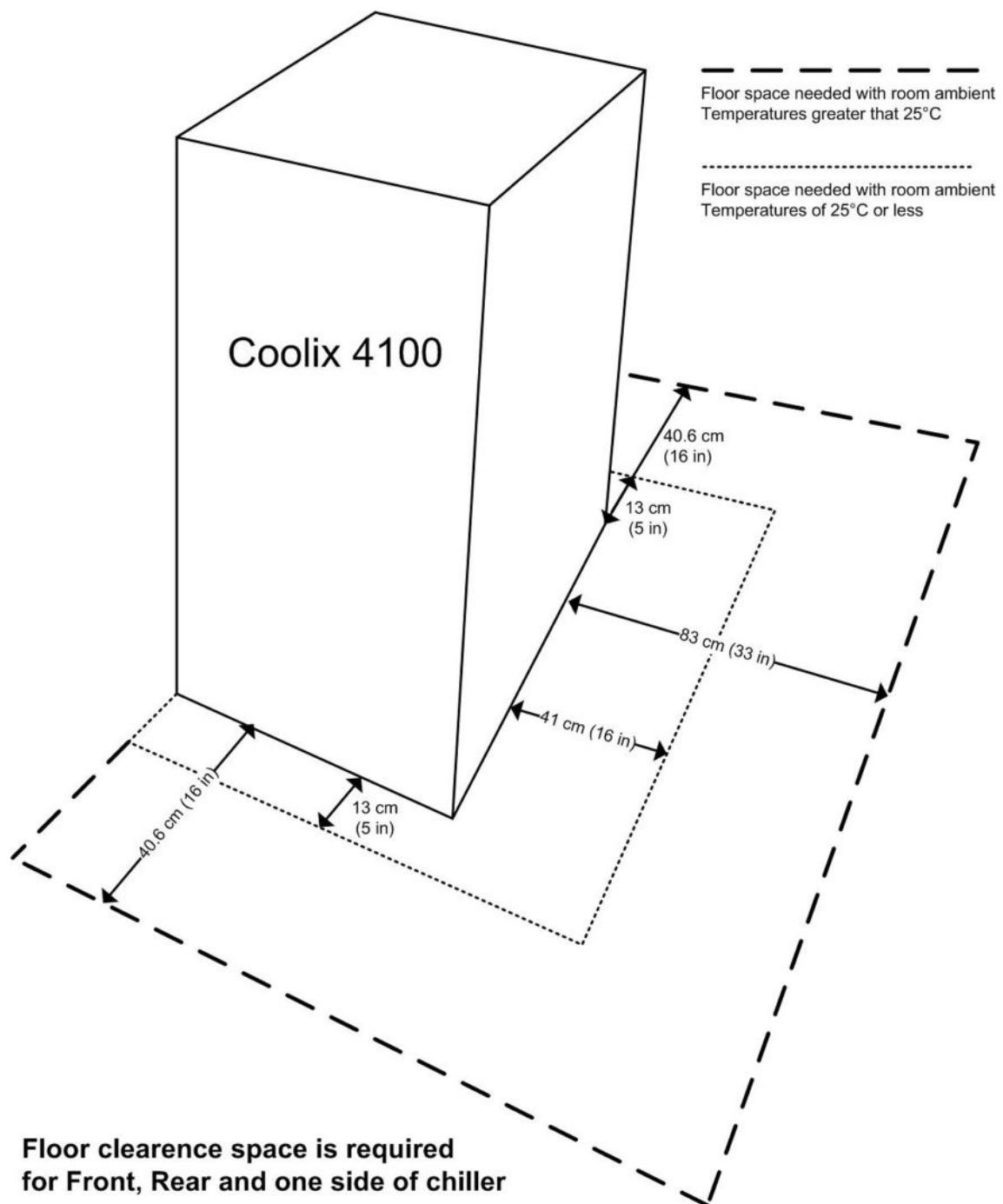
⇒ : AIR FLOW

Illustration 2-36: Dimension Diagram for X-Ray Tube Chiller 4100



NOTE: Required floor space depends on ambient room temperatures. When in doubt, allow for maximum floor space.

Illustration 2-37: Floor Space Diagram for X-Ray Tube Chiller 4100



NOTICE
The chiller cannot be located more than 3 m (10 ft) below the detector

Illustration 2-38: Detector Chiller Thermo-Con Dimensions & Orientation

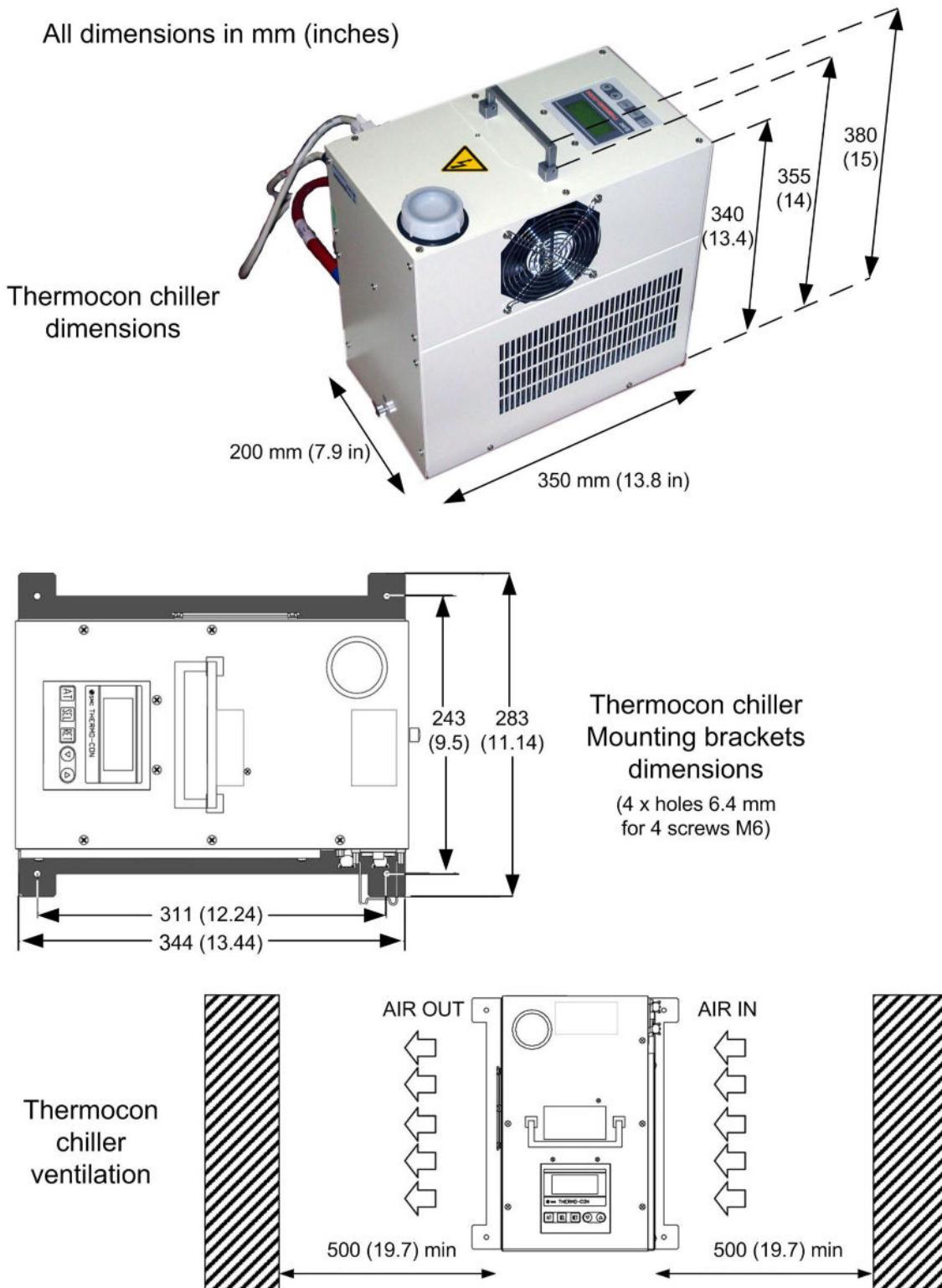
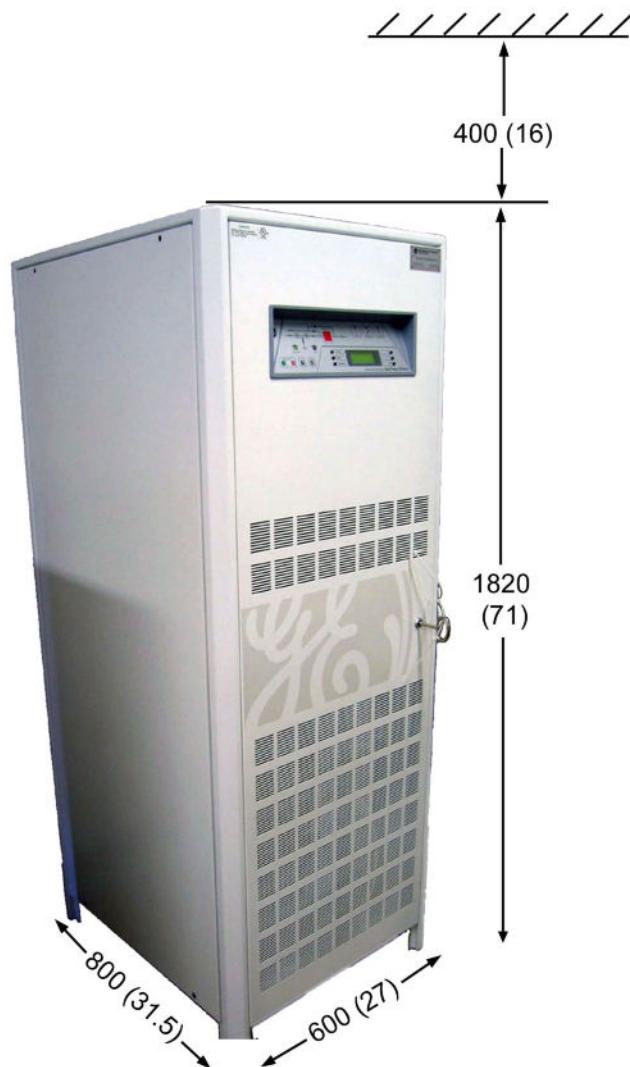


Illustration 2-39: Fluoro UPS UL Layout (Optional)

All dimensions are in mm (inches)



The left, right or back side of the UPS cabinet can be positioned against the wall.

The front side of the UPS cabinet must be accessible for maintenance operation.

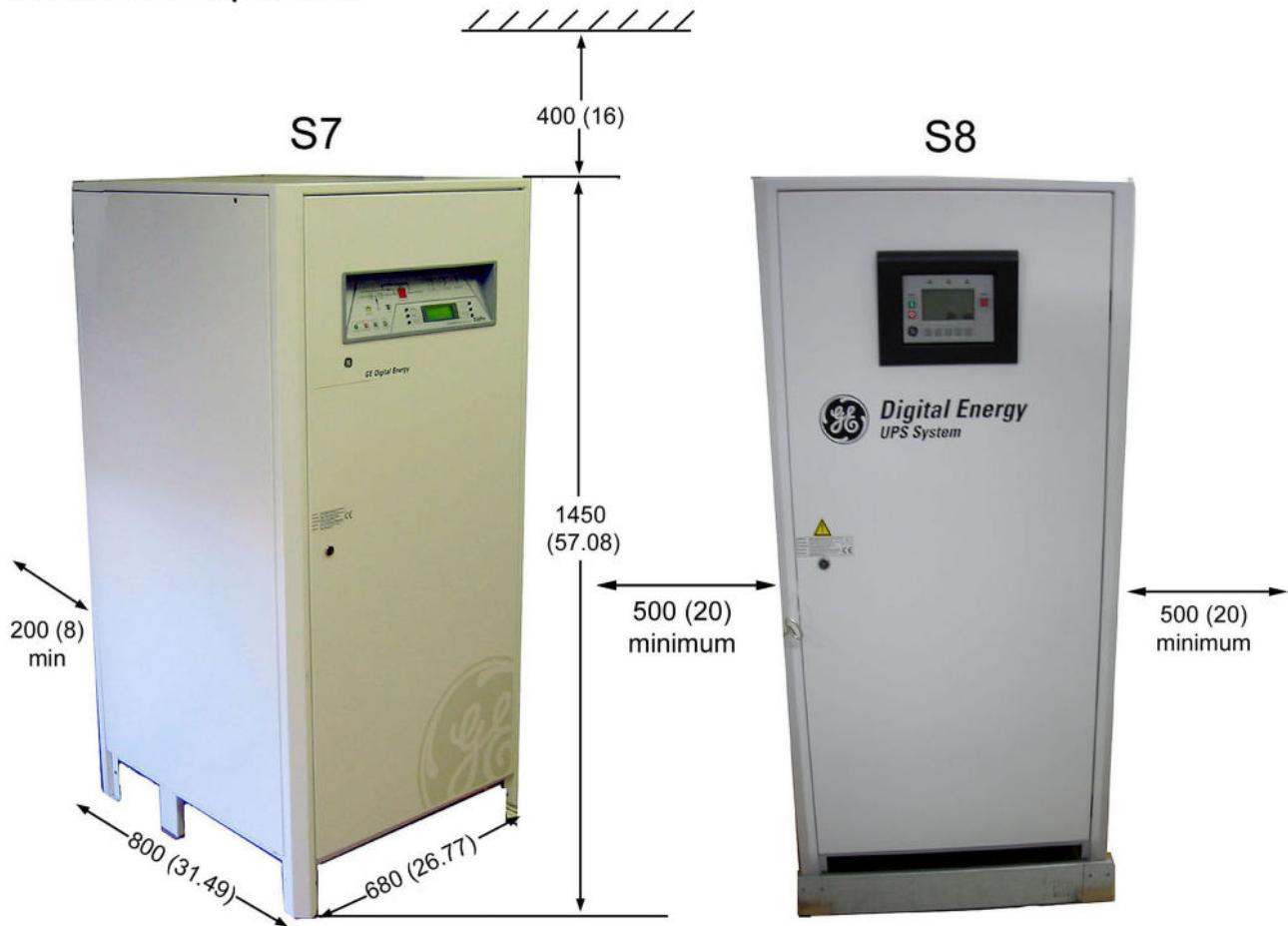
In front of the cabinet, the clear width of the service area to insure electrical safety shall be at least 0.9 m. In cases where 2 cabinets are installed face to face (both sides of the access way), the clear width shall be at least 1.2 m.

Recommended minimum clearance between ceiling and top of the UPS should be 400 mm (16") for proper cooling air exhaust.

Illustration 2-40: Fluoro UPS CE Layout (Optional)

Both S7 and S8 series have the same dimensions and clearance requirements

All dimensions are in mm (inches)



To improve the ventilation and to accommodate the maintenance operations for UPS and battery, a minimum distance of 200 mm (8") from the wall is required.

The right and left side of the UPS cabinet must be accessible for maintenance operation.

In front of the cabinet, the clear width of the service area to insure electrical safety shall be at least 0.9 m. In cases where 2 cabinets are installed face to face (both sides of the access way), the clear width shall be at least 1.2 m.

Recommended minimum clearance between ceiling and top of the UPS should be 400 mm (16") for proper cooling air exhaust.

NOTE: A minimum distance of 2 m (79 in) between X-Ray tube Chiller and Fluoro UPS CE cabinet is required.

NOTE: A Fire extinguisher (non-water type, ex. CO²) must be installed close to the Fluoro UPS CE cabinet.

Illustration 2-41: UPS IF Box (Optional)

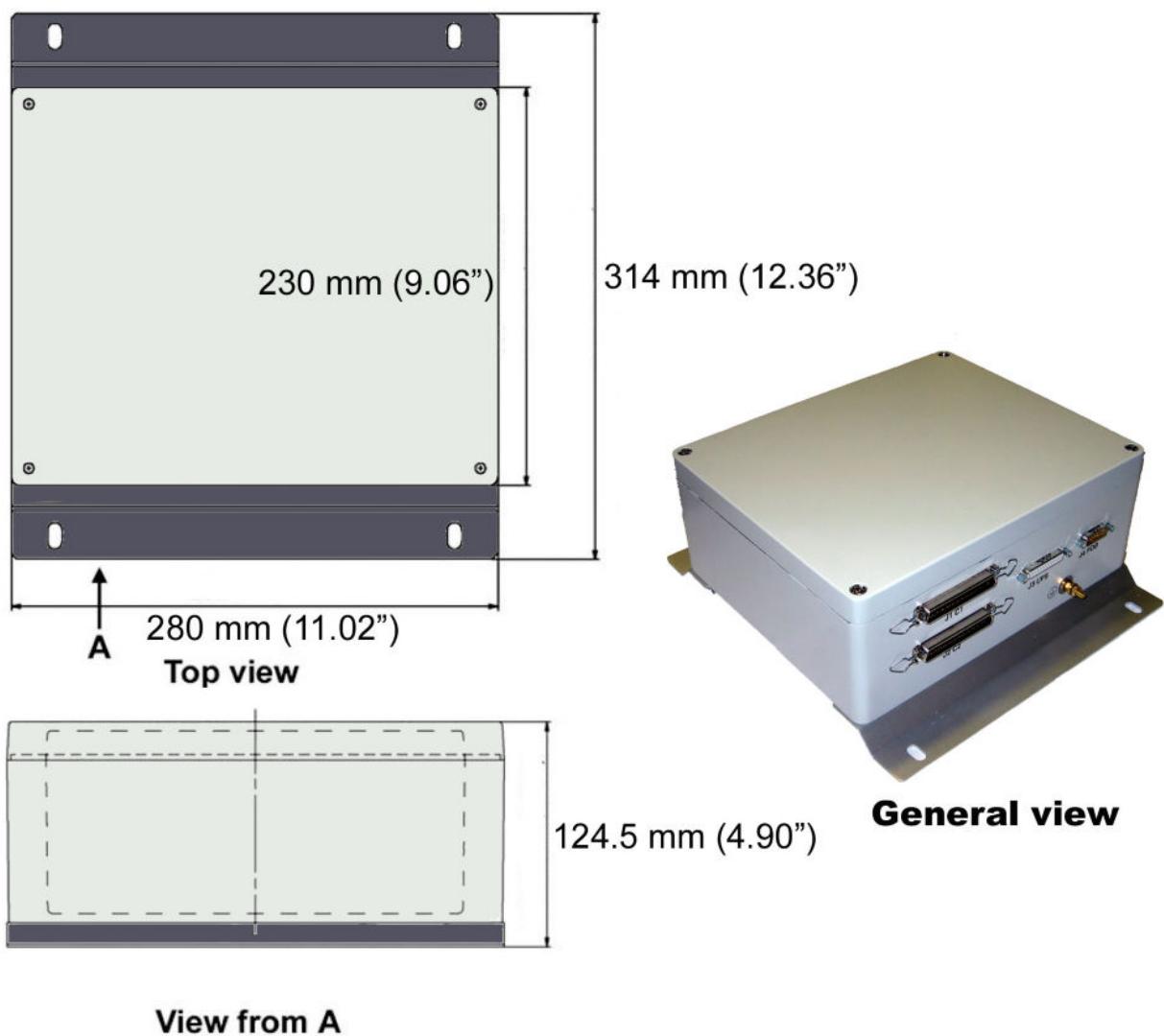


Illustration 2-42: 3 kVA UPS



NOTE: The height of the 3 kVA UPS is 325 mm (12.79 in) without feet.

Illustration 2-43: EMI Filter Box (CE Only)

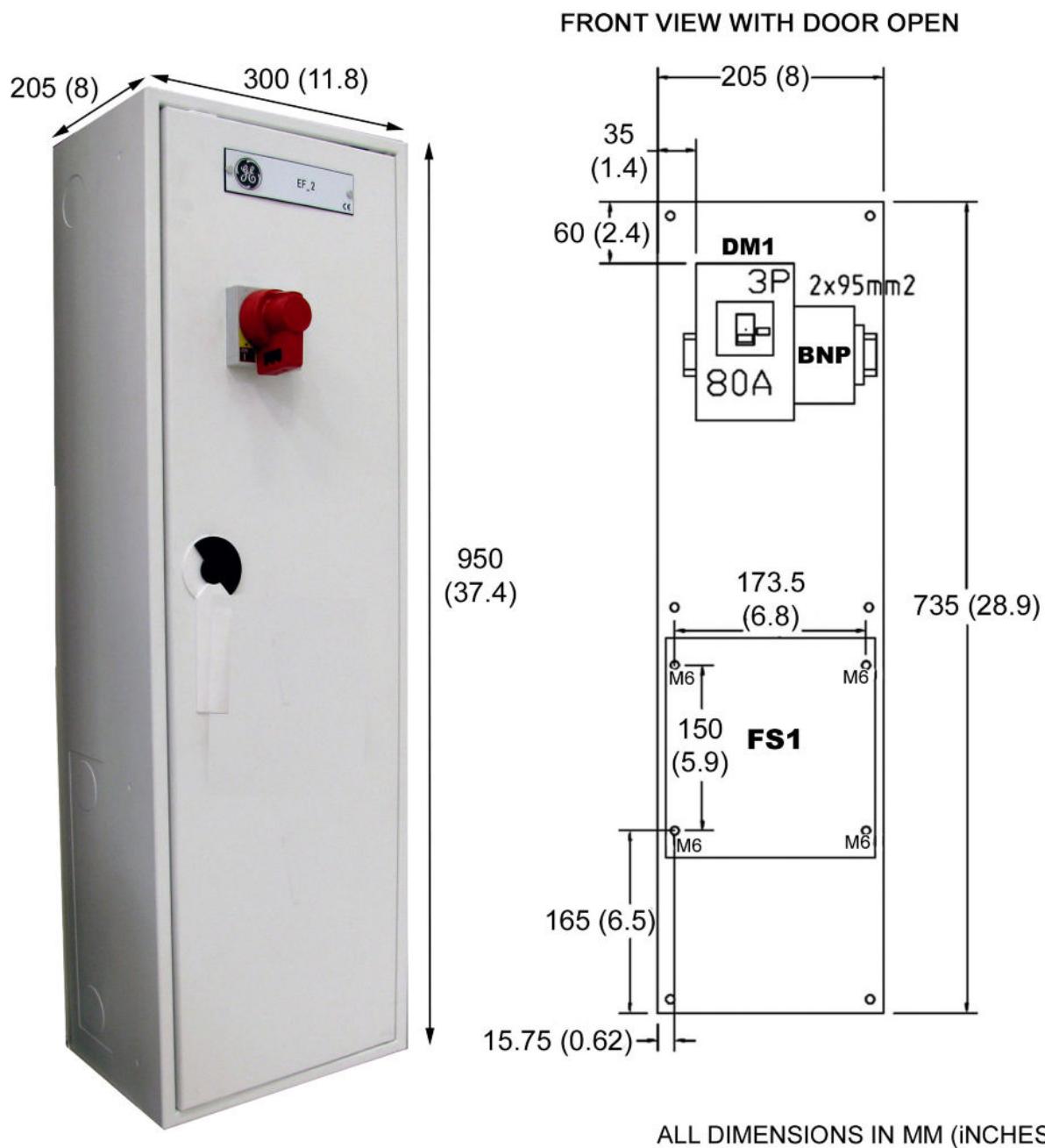
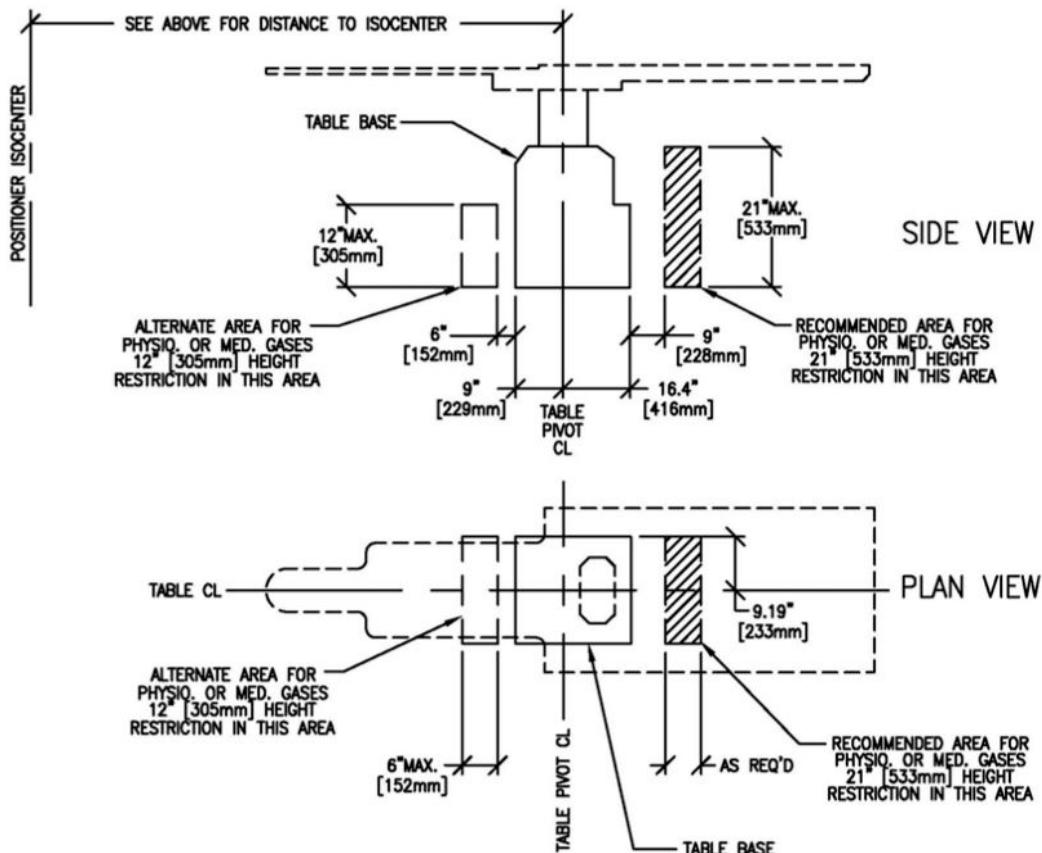


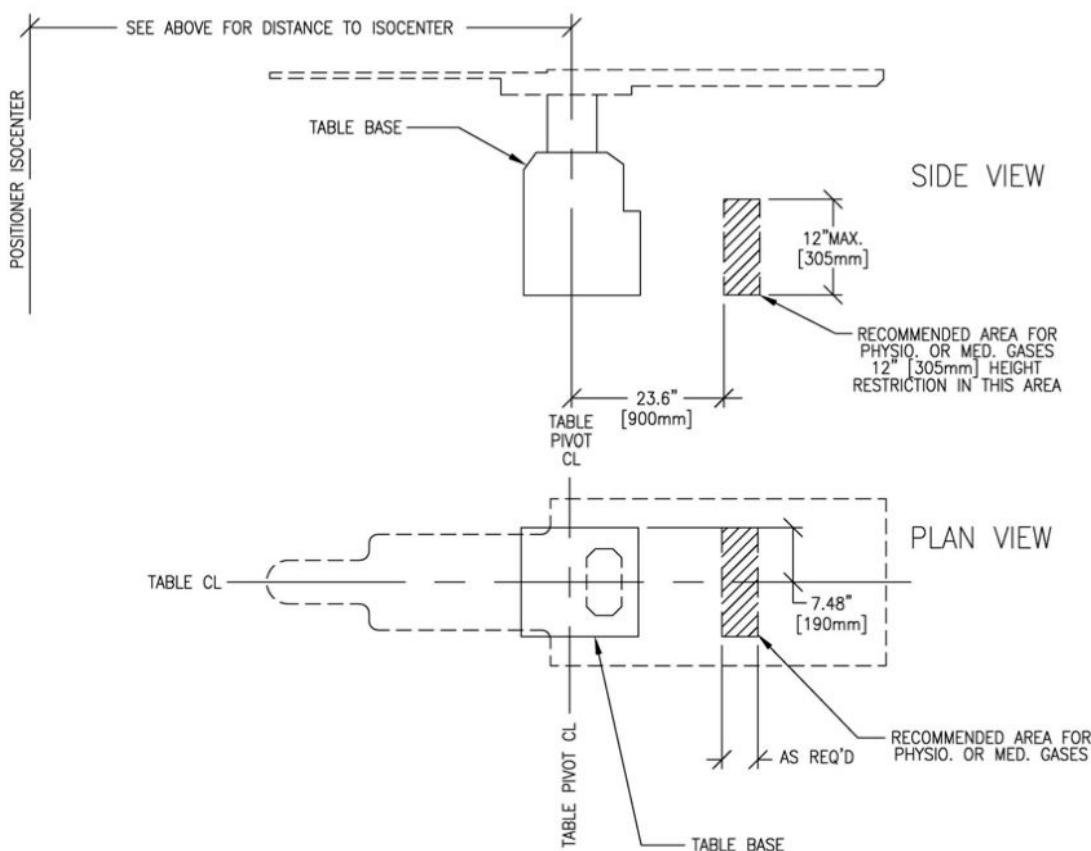
Illustration 2-44: Gas box outlets Omega IV



NOTE: With a Tilting table, the minimum distance from table pivot to the medical Gas Box is 600 mm and the maximum dimensions of the medical Gas Box are :

- height = 305 mm
- width = 250 mm
- length = 500 mm

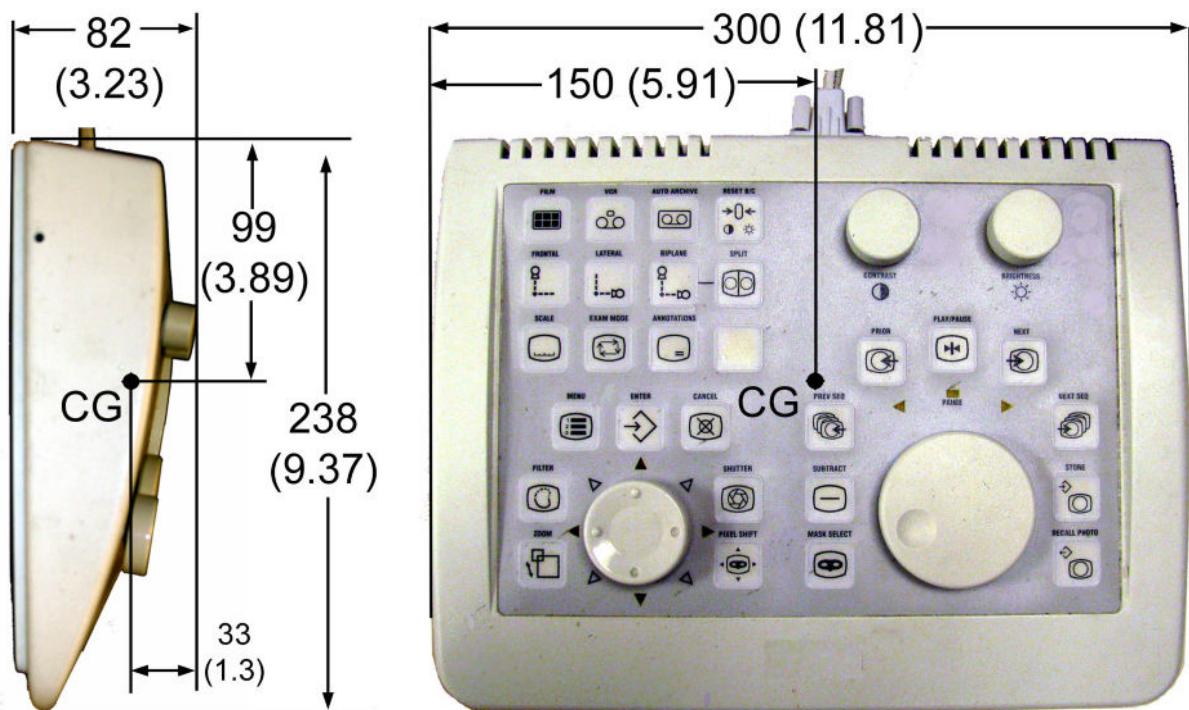
Illustration 2-45: Gas box outlets Omega V



NOTE: With a Tilting table, the minimum distance from table pivot to the medical Gas Box is 600 mm and the maximum dimensions of the medical Gas Box are :

- height = 305 mm
- width = 250 mm
- length = 500 mm

Illustration 2-46: DL Keypad Dimensions



All dimensions are in mm (inches)

Illustration 2-47: DL Image Monitor Dimensions

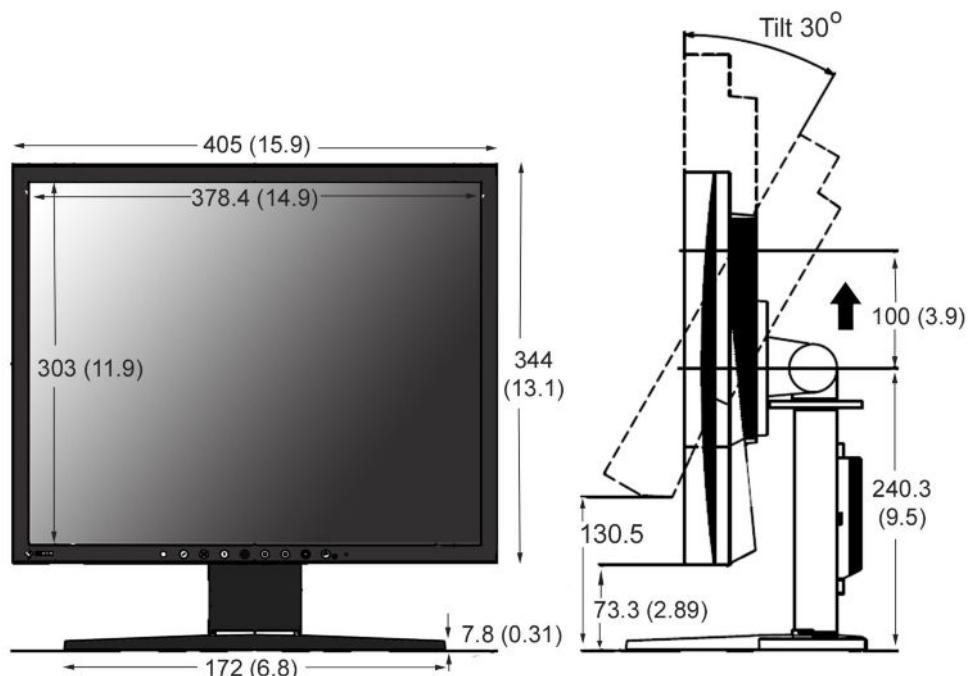
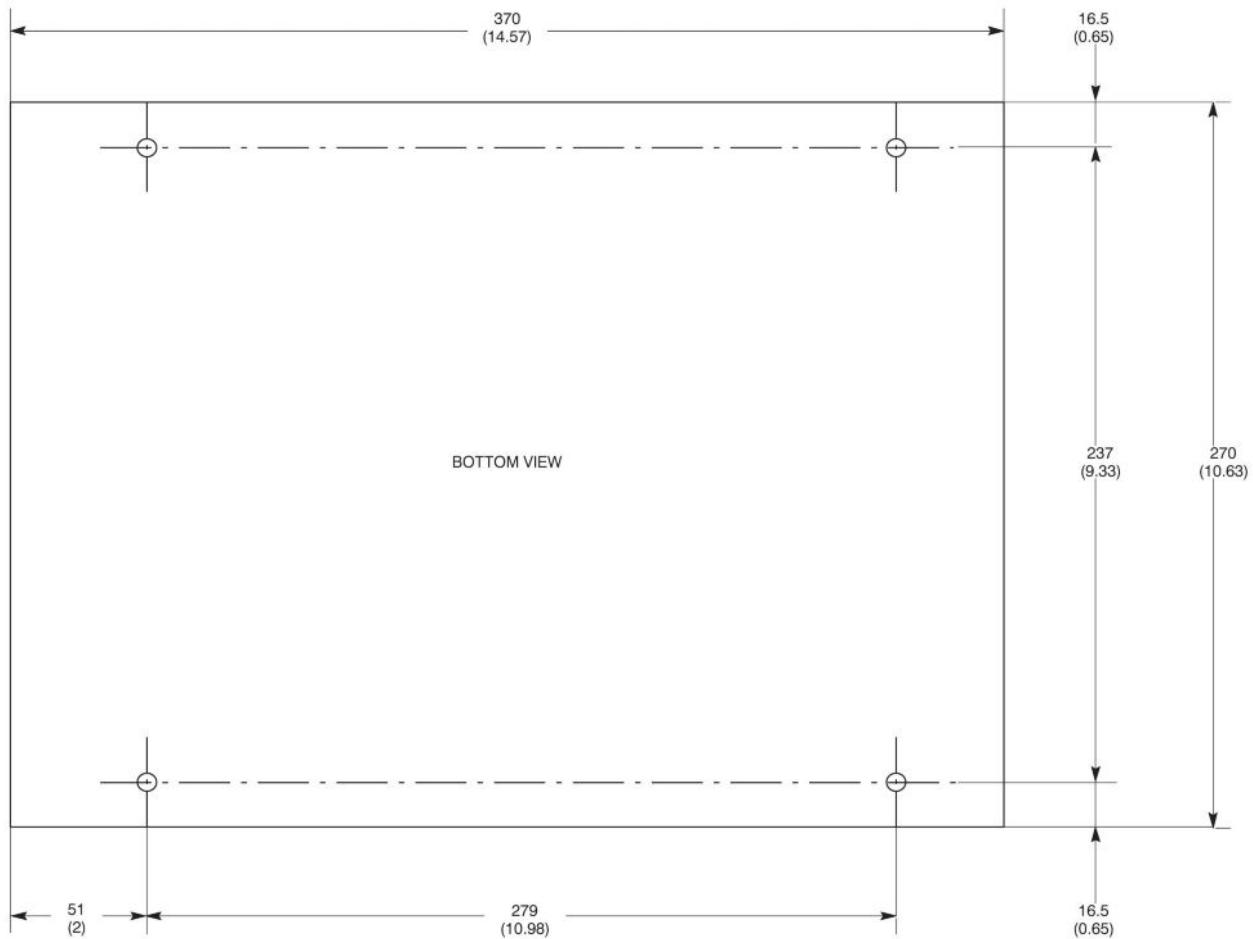


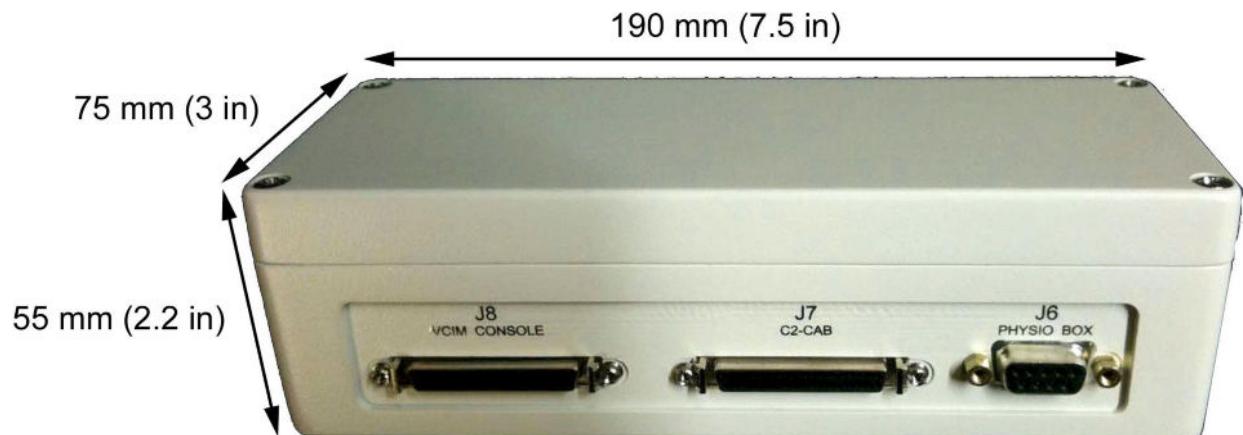
Illustration 2-48: VCR Mounting Holes Location

All dimensions are in mm (inches)

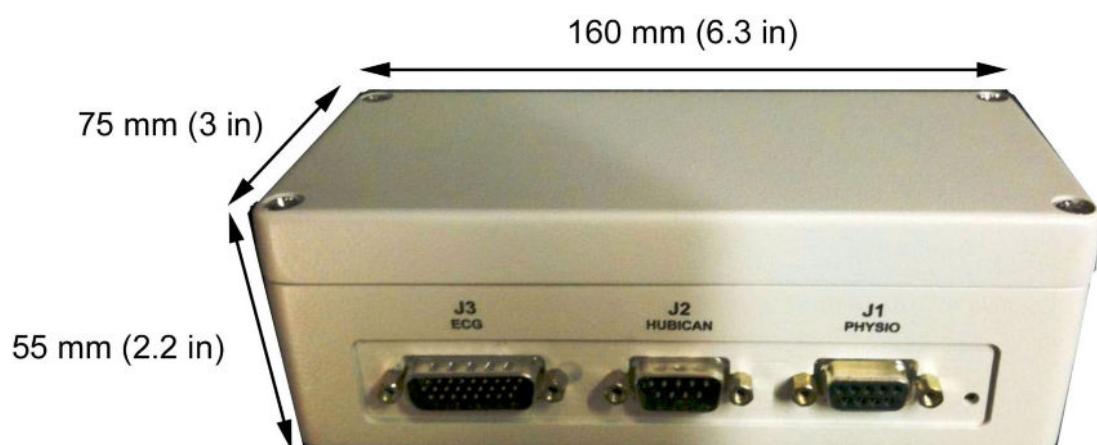


NOTE: Unscrew the four *feet* on the bottom. Use these four holes to fix the VCR. The mounting holes accommodate M3 x 10 mm screws. The holes are 0.4 inches (10 mm) deep.

Illustration 2-49: ECG Acquisition Device Modules



Hubican Module



Physio Module

Illustration 2-50: Large Display cabinet dimensions (Optional)

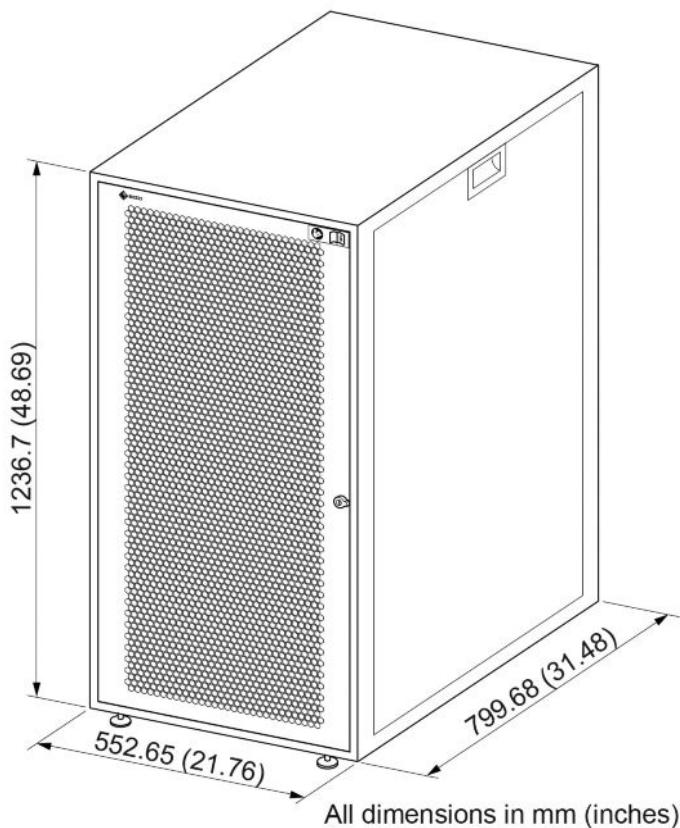
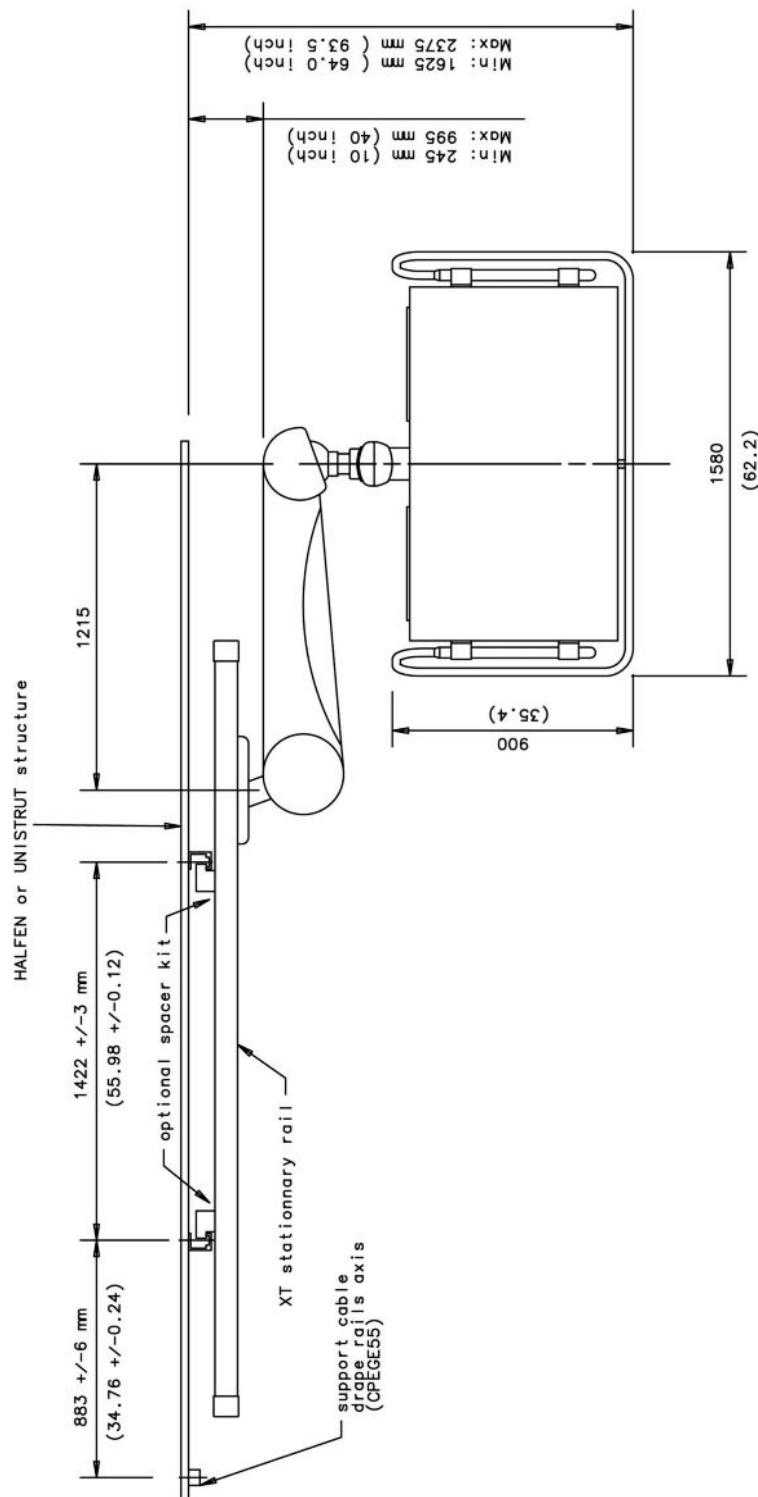


Illustration 2-51: Large Display suspension dimensions (Optional)



1.4 Basic Innova System Compatibility

1.4.1 System Compatibility Cross-Reference - Gantry

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
LC ANGIO POSITIONER	2341727	This document	HV cable 24 m
LC COMBO POSITIONER	2396420	This document	
LC CARDIO POSITIONER	5126671	This document	
C2 CABINET	2335139-2/2335139-4	This document	
SMART BOX	5408840	This document	Omega Table - Silver Edition
SMART HANDLE	5408842	This document	Omega Table - Silver Edition
SMART BOX	5413420	This document	Innova ^{IQ} Table - Silver Edition
SMART BOX IPX4	5407800	This document	OR Table - Silver Edition
BOLUS HANDLE	2111431	This document	Optional

1.4.2 System Compatibilities Cross-Reference - Tables Sub-System

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
OR TABLE	Table without covers	5142213-001	This document
	OR Table Top Telescopic Covers	5413797	
INNOVA ^{IQ} TABLE	Table without covers	5142213-001	This document
	Innova ^{IQ} Table Top Telescopic Covers	5413797-2	
OMEGA IV TABLE	2320221-3	This document	Includes tabletop
OMEGA V LONG TABLE	2320045-5	This document	Includes tabletop
OMEGA V NON MOTORIZED LONG TABLE	2320045-6	This document	Includes tabletop
TSSC	5408841	This document	Omega Table - CDRH Silver Edition
TSSC	5413423	This document	Innova ^{IQ} Table - CDRH Silver Edition
TSSC IPX4	5407798	This document	OR Table - CDRH Silver Edition
TABLE PANNING DEVICE IPX4	5184673-2	NA	

1.4.3 System Compatibilities Cross-Reference - Jedi X-Ray Generator

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
JEDI 100 VASC	2326480	NA	Includes C1 cabinet

1.4.4 System Compatibility Cross-Reference - X-Ray Head

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
Performix 160 A X-Ray tube	D2801A	Not applicable	

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
Cardio collimator	5127609-2	Not applicable	
Angio collimator	5245319	Not applicable	
COOLIX 4100 Chiller	5416114	Not applicable	
Coolix 4100 Autotransformer	5416111 (UL version) or 5416110 (IEC version)	Not applicable	

1.4.5 System Compatibility Cross-Reference - Innova System Imaging And X-Ray Control

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	NOTES
VCIM	2340519	NA	
DL Liberty user interface	2345576	This document	
C1 cabinet	2335129-8	This document	
Innova IGS 520, 21 cm Revolution Digital detector	5127984	This document	
Innova IGS 530, 31 cm Revolution Digital detector	2359035	This document	
Innova IGS 540, 41 cm Revolution Digital detector	2329766	This document	
Detector Chiller Thermo-Con	5341164	This document	

1.4.6 System Compatibility Cross-Reference - Monitor Support & Suspension

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
PRECABLED LCD 4 MONITOR SUSPENSION: • CABLE HARNESS 24 m • or CABLE HARNESS 36 m	5126894 • 2378538 • or 2378540	2393190-100	
PRECABLED LCD 6 MONITOR SUSPENSION: • CABLE HARNESS 24 m • or CABLE HARNESS 36 m	5126896 • 2378538 • or 2378540	2393190-100	

1.4.7 System Compatibilities Cross-Reference 19" Monitors

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
LCD 19" - Eizo SMD19100G B&W with Stand	5148721-2		
LCD 19" - Eizo SMD19100G B&W without Stand	5148721-3		
LCD 19" - Eizo SMD19100G Color with Stand	5148720-2	sm 5219983-100	
LCD 19" - Eizo SMD19100G Color without Stand	5148720-3		

Eizo 19" LCD HB color monitor RX150 GE		sm 5499528-1-8EN	
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1.5 Optional Innova System Components

1.5.1 System Compatibilities Cross-Reference - Omega Table Sub-System

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
SMART HANDLE	5408842	This document	
SMART BOX	5408840	This document	
TSSC	5408841	This document	

1.5.2 System Compatibilities Cross-Reference - Innova^{IQ} Table Sub-System

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
SMART BOX	5413420	This document	
TSSC	5413423	This document	

1.5.3 System Compatibilities Cross-Reference - OR Table Sub-System

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
SMART BOX	5407800	This document	
TSSC	5407798	This document	

1.5.4 System Compatibilities Cross-Reference - Wall Mount Monitor

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
Wall Mount for 17 or 21" (43 or 53 cm)	2305202	2129003-100	
Note: The wall mount monitor is for Control Room only.			

1.5.5 System Compatibilities Cross-Reference - ECG Aquisition

PRODUCT NAME	MODEL NUMBER	PRE-INSTALLATION DOCUMENT NUMBER	CATALOG NUMBER
ECG Acquisition Kit		This document	

2 Room Layouts

2.1 Room Dimension Requirements



WARNING

LOCATION IN TECHNICAL ROOM FOR ELECTRICAL CABINETS IS MANDATORY.

THE ELECTRONIC CABINETS (C1, C2, OPTIONAL LD CABINET, COOLIX 4100 CHILLER, DETECTOR CHILLER AND FLUORO UPS WHEN INSTALLED) INCLUDE FANS THAT ARE CREATING AIR-CIRCULATION OF PULSED-AIR. WHEN THIS PULSED AIR IS IN AN ENVIRONMENT THAT MAY CONTAIN AIRBORNE PATHOGENS LIKE AN EXAM ROOM/CONTROL ROOM, THERE IS A RISK OF TRANSMISSION OF THESE AIRBORNE PATHOGENS FROM PATIENTS TO OTHER PATIENTS OR CLINICAL PERSONNEL (NOSOCOMIAL DISEASES).

TO REDUCE THIS RISK, THE ELECTRONIC CABINETS MUST BE INSTALLED IN A ROOM SEPARATED FROM EXAM ROOM/CONTROL ROOM, I.E., TECHNICAL ROOM.

For exam and control rooms, several configurations are recommended.

NOTE: Motion controls installed in remote location from the table shall be installed at a location where all the positioner axis are visible by the operator.

In case the system has a tilting table, the remote motion controls shall not be installed on the longitudinal axis of the Table (to avoid any operator visual dead angle due to tilted table top hiding the patient).

Refer to [Patient Room Layout](#) and [Control Room Layout](#) to see recommended exam / control rooms layouts.

Table 2-2: Exam Room dimensions

	Length	Width	Ceiling Height
Recommended:	32 ft 0 in (9.75m)	20 ft 0 in (6.00 m)	10 ft 0 in (3.05 m)
Minimum:	For minimum exam room lengths, see different table configurations Table 2-3 , Table 2-4 , Table 2-5 , Table 2-6 and Illustration 2-52	14 ft 5 in (4.40 m)	9 ft 0 in (2.74 m)

For the tables below, refer to [Illustration 2-52](#)

Note: For Head Extender Dimensional drawing, refer to [Dimension Drawings](#)

Table 2-3: Omega IV Table

Configuration	Minimum Room length in mm (inches)	
	without Head Extender (see note above)	with Head Extender (see note above)
Cardio 1395 mm (54.9 in)	5470 (215.3)	5470 (215.3)

Table 2-4: Omega V Table

Configuration	Minimum Room length in mm (inches) A	
	without Head Extender	with Head Extender F
Angio 1278 mm (50.3 in) C	6036 (237.6)	6036 (237.6) + 150 (6)
Cardio and Neuro 1395 mm (54.9 in) B	6036 (237.6)	6036 (237.6) + 150 (6)

Table 2-5: Tilting table

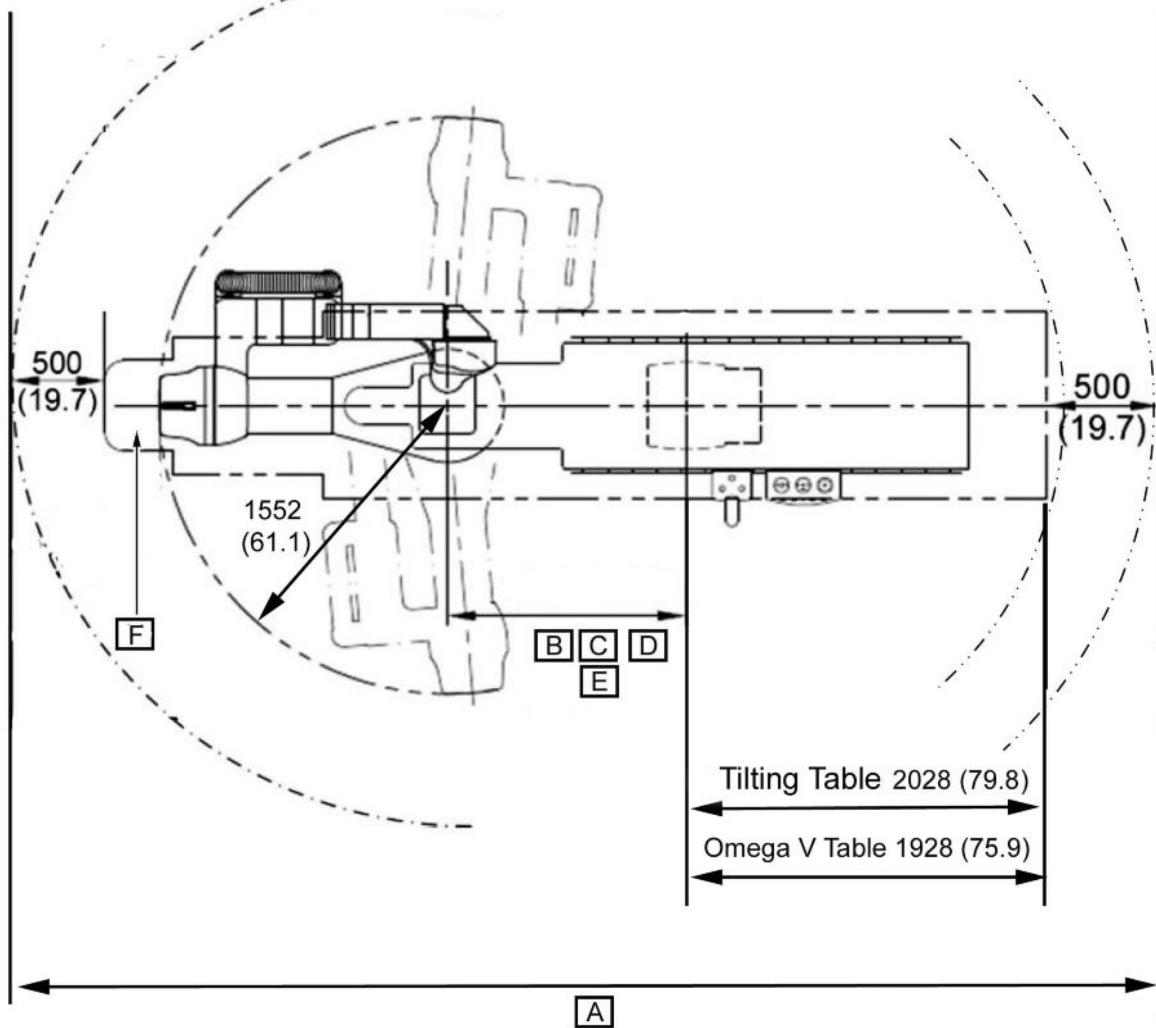
Configuration	Minimum Room length in mm (inches) A	
	without Head Extender	with Head Extender F
Angio 1278 mm (50.3 in) C	6136 (241.6)	6136 (241.6) + 150 (6)
Neuro 1395 mm (54.9 in) D	6136 (241.6)	6136 (241.6) + 150 (6)

Table 2-6: Tilting table Upgrade

Configuration	Minimum Room length in mm (inches) A	
	without Head Extender	with Head Extender F
Angio 1278 mm (50.3 in) C	6136 (241.6)	6136 (241.6) + 150 (6)
Neuro 1395 mm (54.9 in) D	6136 (241.6)	6136 (241.6) + 150 (6)
Neuro (old distance, no longer used) 1595 mm (62.8 in) E	6175 (243.1)	6175 (243.1) + 150 (6)

Illustration 2-52: Minimum Room Length Dimension

All dimensions are in mm (inches)



2.2 Room Layout Drawings

2.2.1 Patient Room Layout

2.2.1.1 Room Layout for Innova System (examples of room layouts)



WARNING

LOCATION IN TECHNICAL ROOM FOR ELECTRICAL CABINETS IS MANDATORY.

THE ELECTRONIC CABINETS (C1, C2, OPTIONAL LD CABINET, COOLIX 4100 CHILLER, DETECTOR CHILLER AND FLUORO UPS WHEN INSTALLED) INCLUDE FANS THAT ARE CREATING AIR-CIRCULATION OF PULSED-AIR. WHEN THIS PULSED AIR IS IN AN ENVIRONMENT THAT MAY CONTAIN AIRBORNE PATHOGENS LIKE AN EXAM ROOM/CONTROL ROOM, THERE IS A RISK OF TRANSMISSION OF THESE AIRBORNE PATHOGENS FROM PATIENTS TO OTHER PATIENTS OR CLINICAL PERSONNEL (NOSOCOMIAL DISEASES).

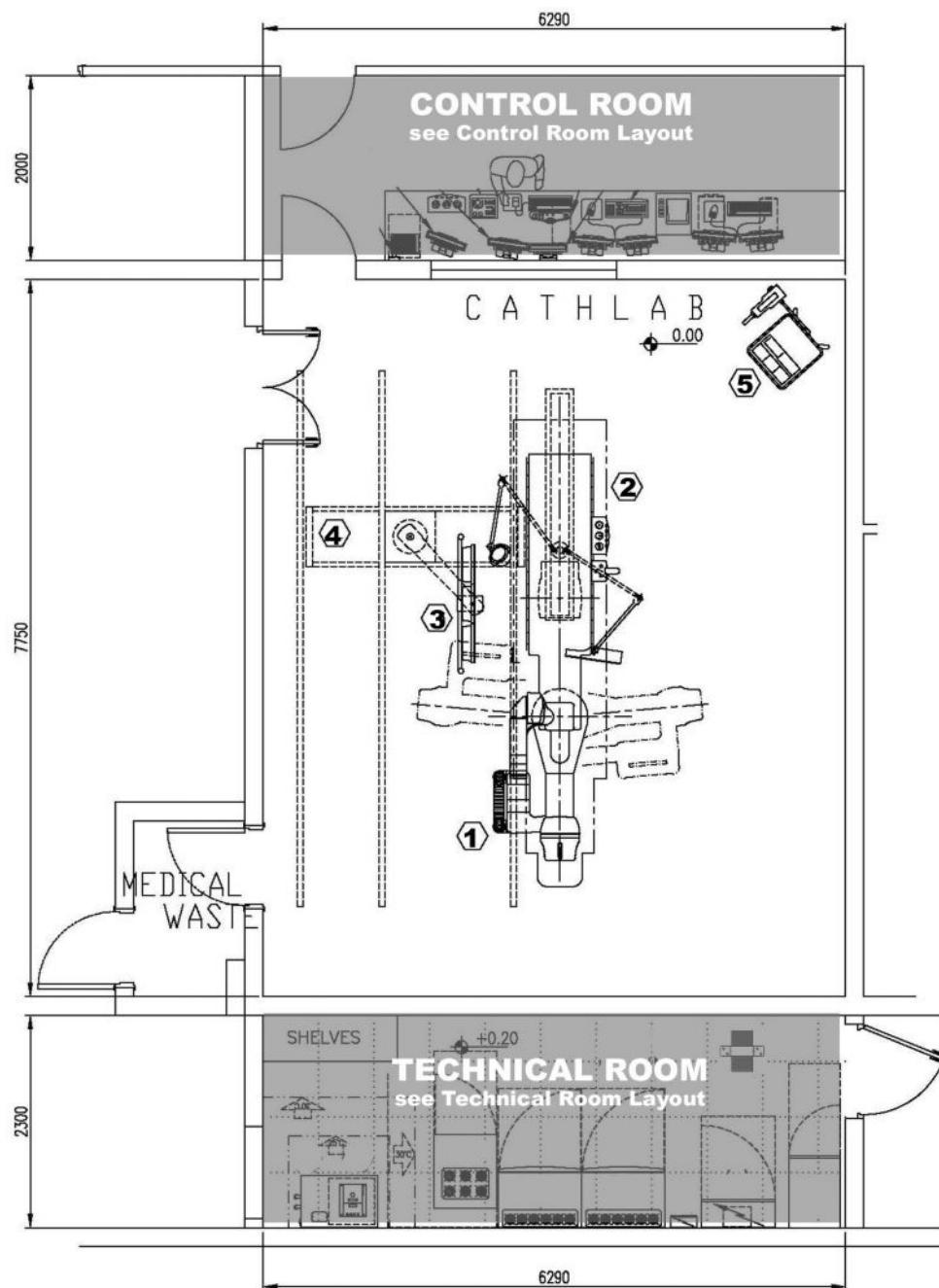
TO REDUCE THIS RISK, THE ELECTRONIC CABINETS MUST BE INSTALLED IN A ROOM SEPARATED FROM EXAM ROOM/CONTROL ROOM, I.E., TECHNICAL ROOM.

Legend of [Illustration 2-53](#), [Illustration 2-54](#), [Illustration 2-55](#) and [Illustration 2-56](#).

1. Gantry
2. Patient table
3. XT monitor suspension
4. Monitor frame
5. Injector on Pedestal

NOTE: The phone outlet must be located less than 1 meter (3 feet) from the C1 Cabinet (3).

Illustration 2-53: Monitor suspensions rails parallel to patient table (to side of table)



WARNING

ILLUSTRATION 2-53 IS MANDATORY WITH HOSPITAL HAVING AIR FLOW
INSTALLATION

Illustration 2-54: Monitor suspensions rails parallel to patient table (over table)

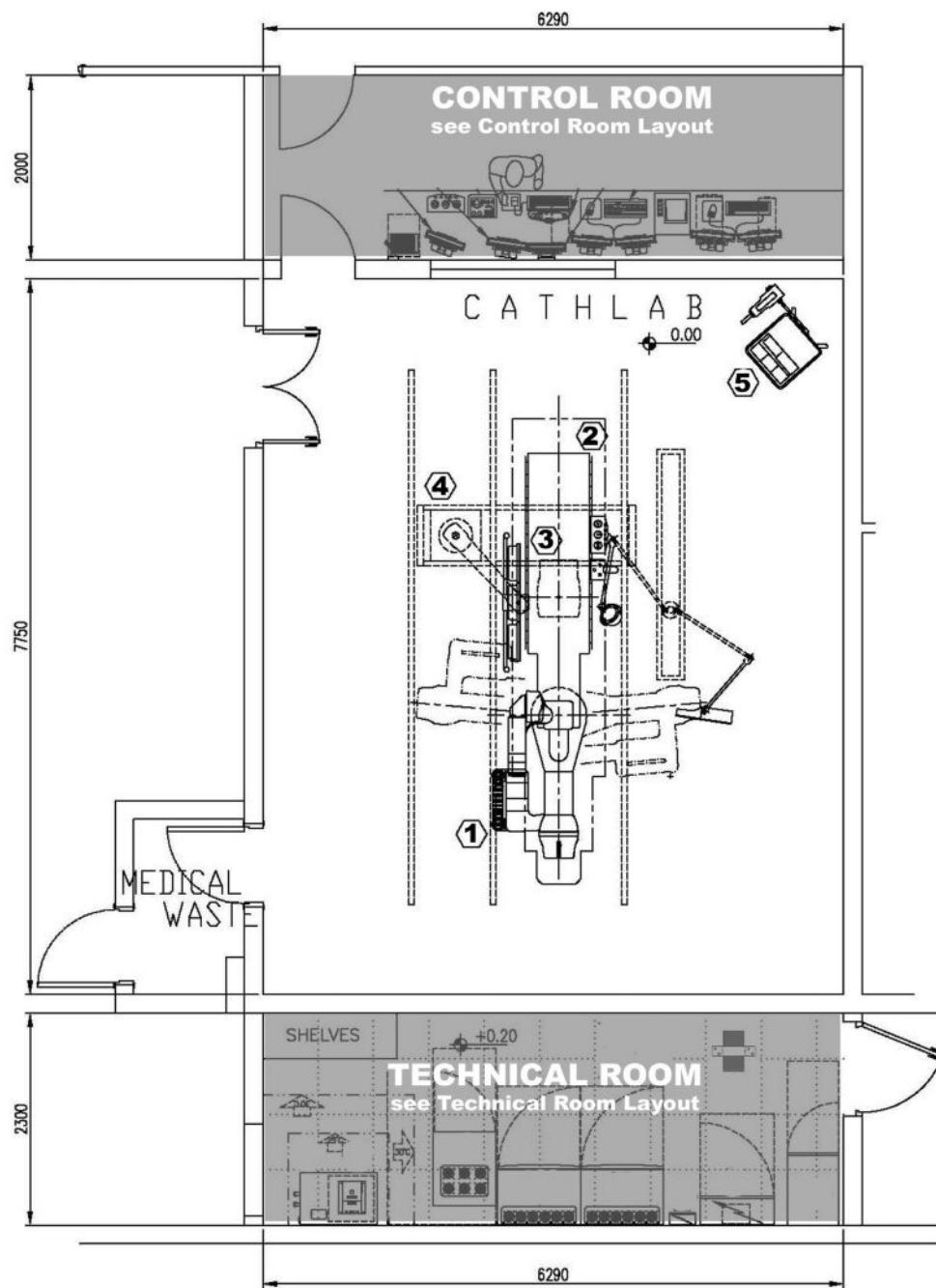


Illustration 2-55: Monitor suspensions rails perpendicular to patient table (Gantry side)

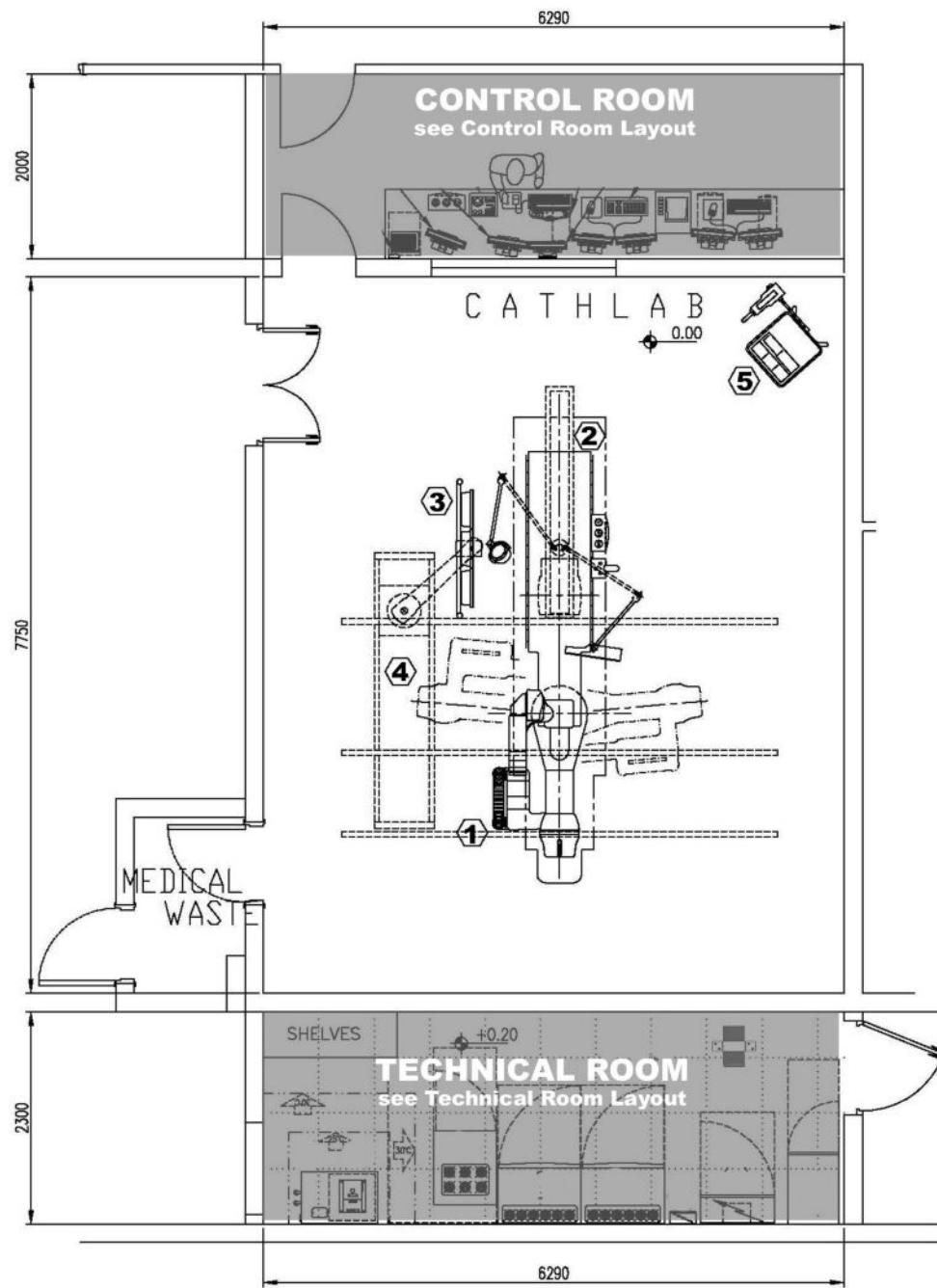
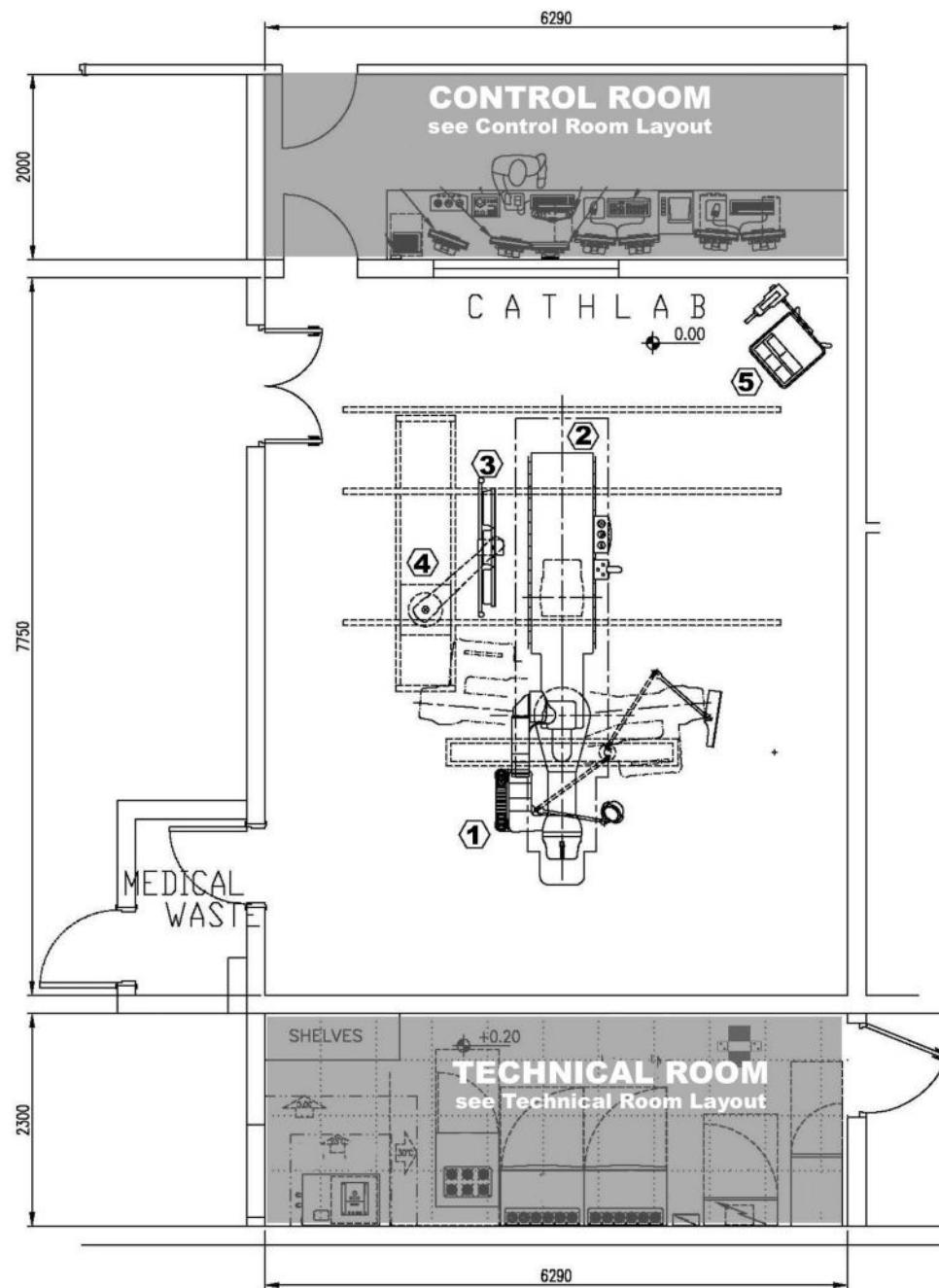
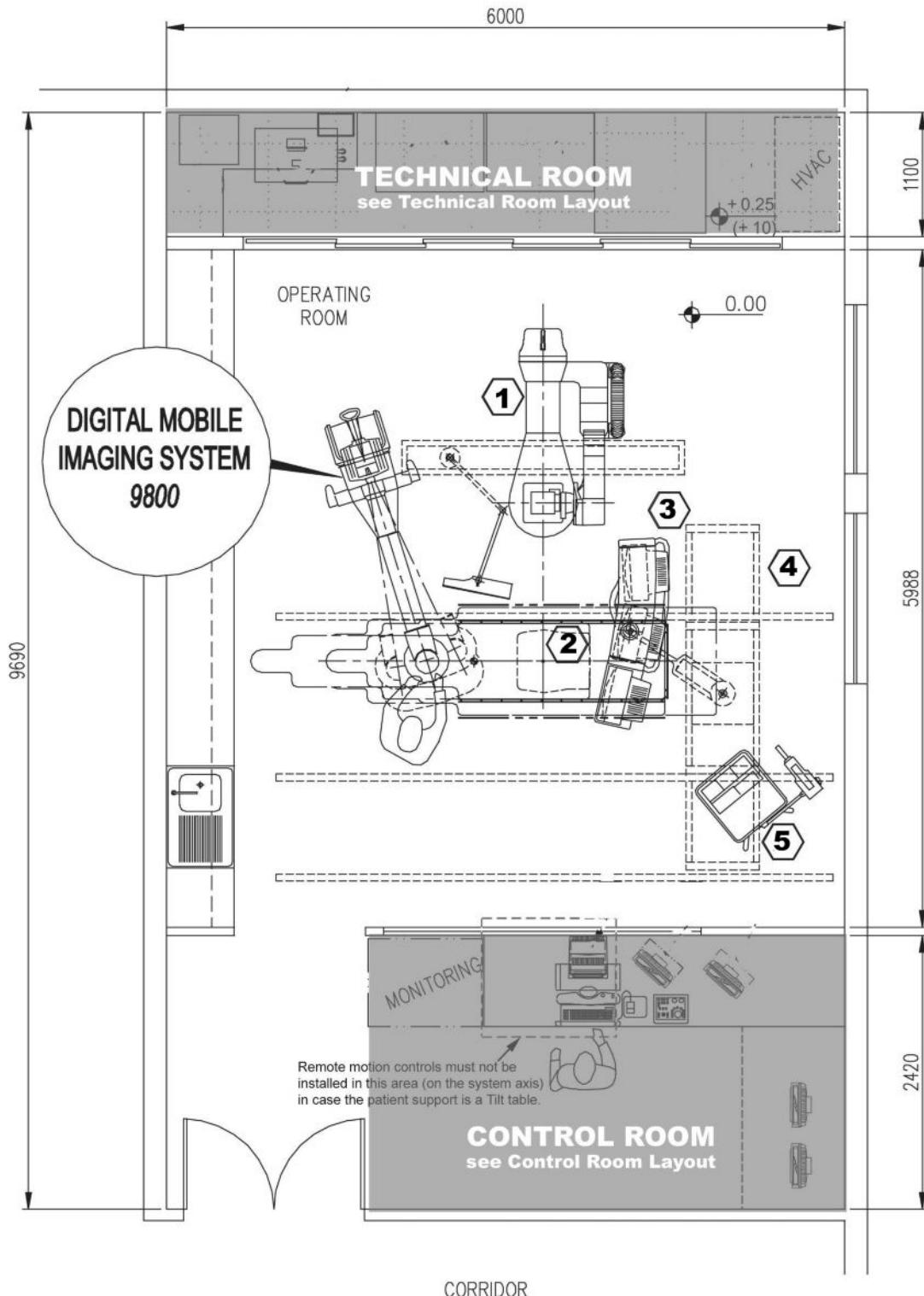


Illustration 2-56: Monitor suspensions rails perpendicular to patient table (table side)



2.2.1.2 Room Layout for Innova System (example with Mobile system 9800)

Illustration 2-57: Room Layout for Innova System



Legend of [Illustration 2-57](#)

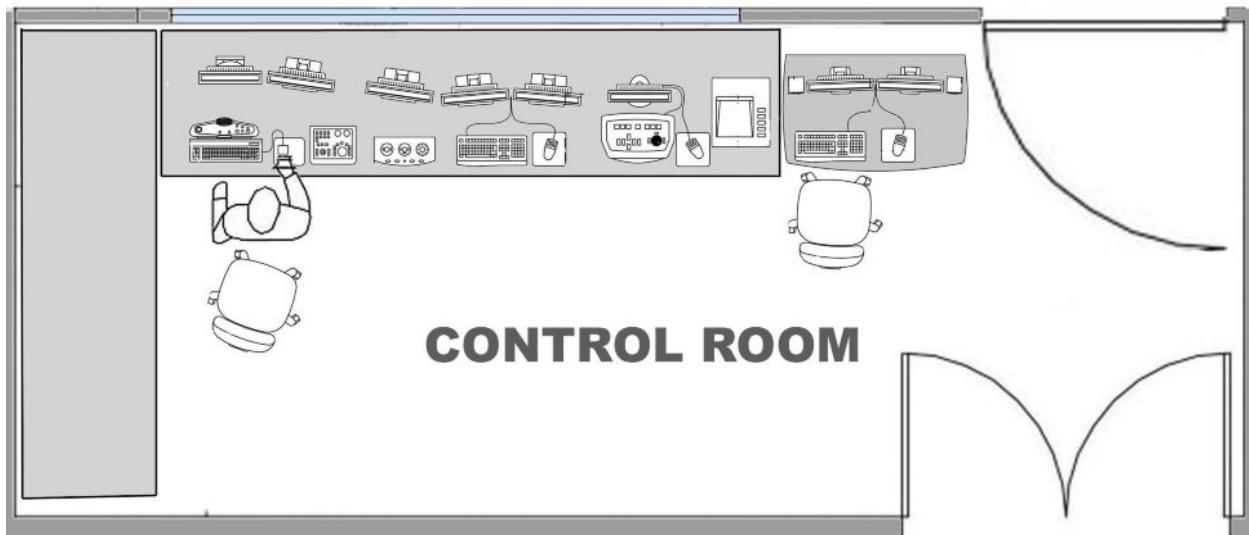
1. Gantry
2. Patient table
3. XT monitor suspension
4. 21" (53 cm) FFD Monitors put on frame with monitoring screen
5. Injector on Pedestal

NOTE: The phone outlet must be located less than 1 meter (3 feet) from the C1 Cabinet (3).

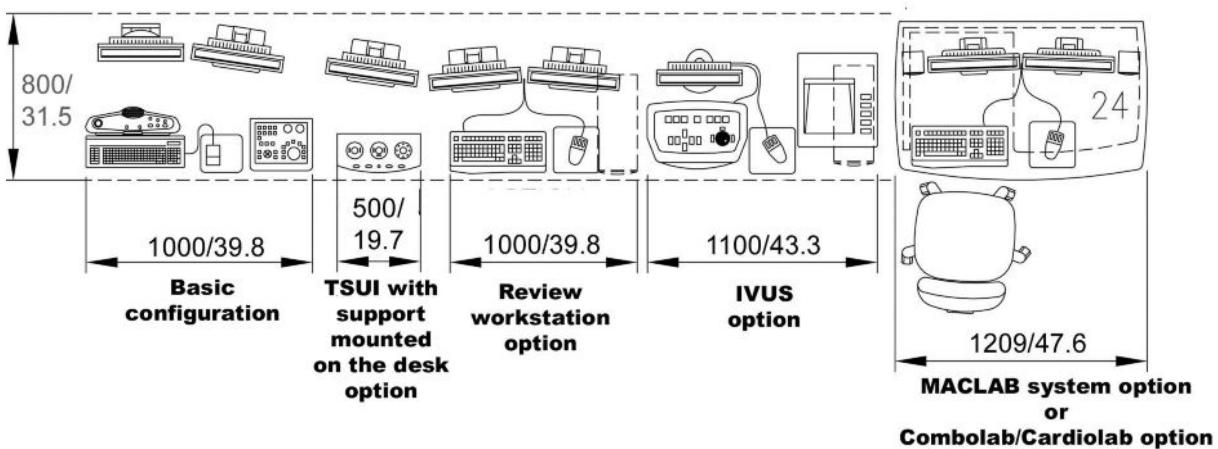
2.2.2 Control Room Layout

Illustration 2-58:

PATIENT ROOM



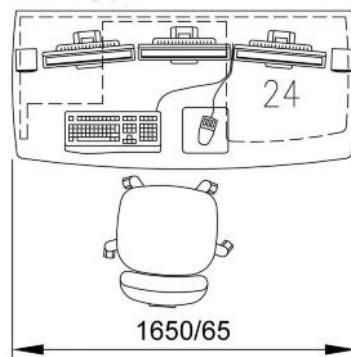
CONTROL ROOM



WARNING

if remote Table User Interface option installed,
ensure full visibility over the moving parts in the
Exam Room (Gantry and Table)

All measurements are in mm (inches)

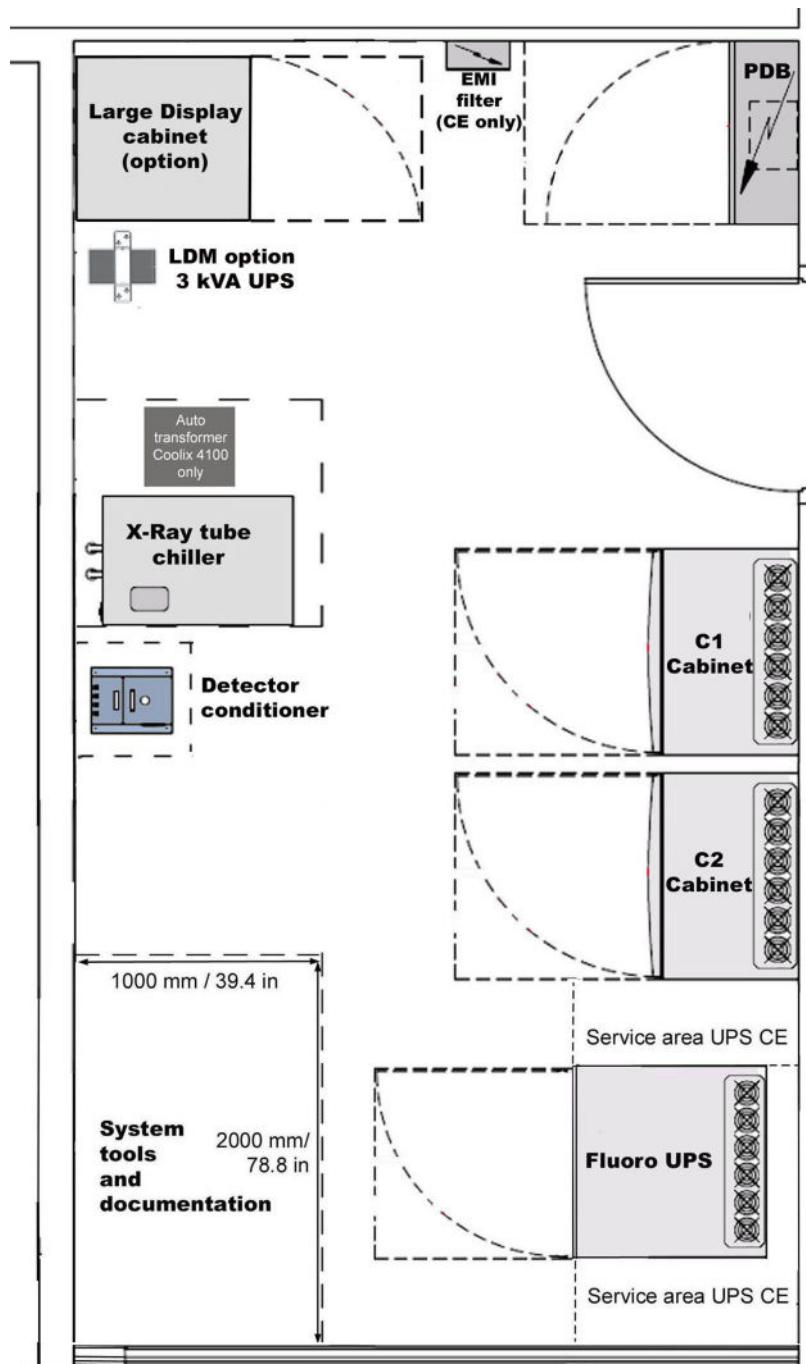


2.2.3 Technical Room Layout

For the service access and ventilation restraint dimensions below, see [Dimension Drawings](#) and [Room Layout Considerations](#).

NOTE: The phone outlet must be located less than 1 meter (3 feet) from the C1 Cabinet.

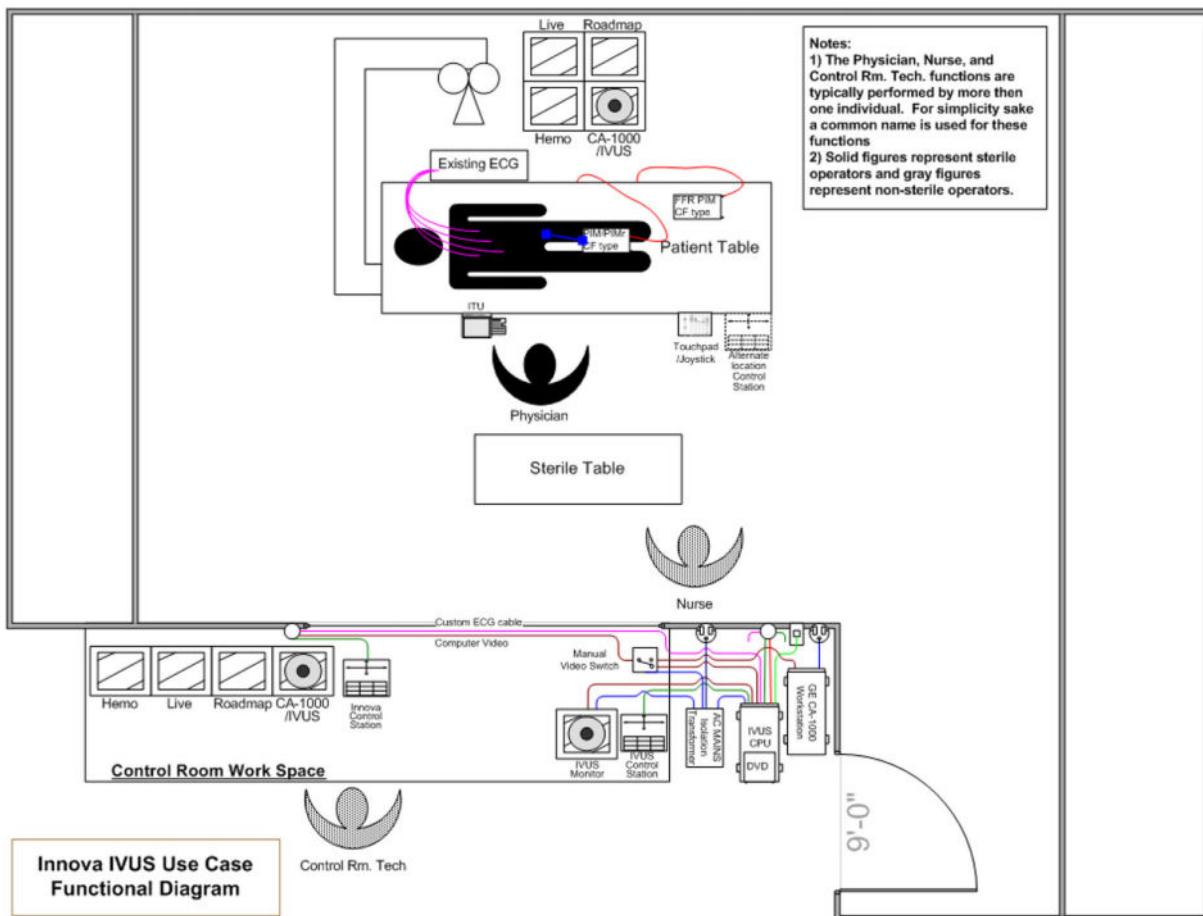
Illustration 2-59:



2.2.4 IVUS Room Layout

2.2.4.1 IVUS Rev 2 and Rev 2+

Illustration 2-60:



NOTE: (For OR Table) the CONTROL STATION shall be installed on a cart.

2.2.4.2 IVUS Rev 3

Refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

2.2.5 ECG Device Room Configurations

The hardware delivered with the ECG acquisition kit will need to be installed in the Control Room and in Exam Room depending on the type of ECG Device used on the site.

The ECG Device/Room configurations need to be checked as this can impact ON the way parts and cables can be installed.

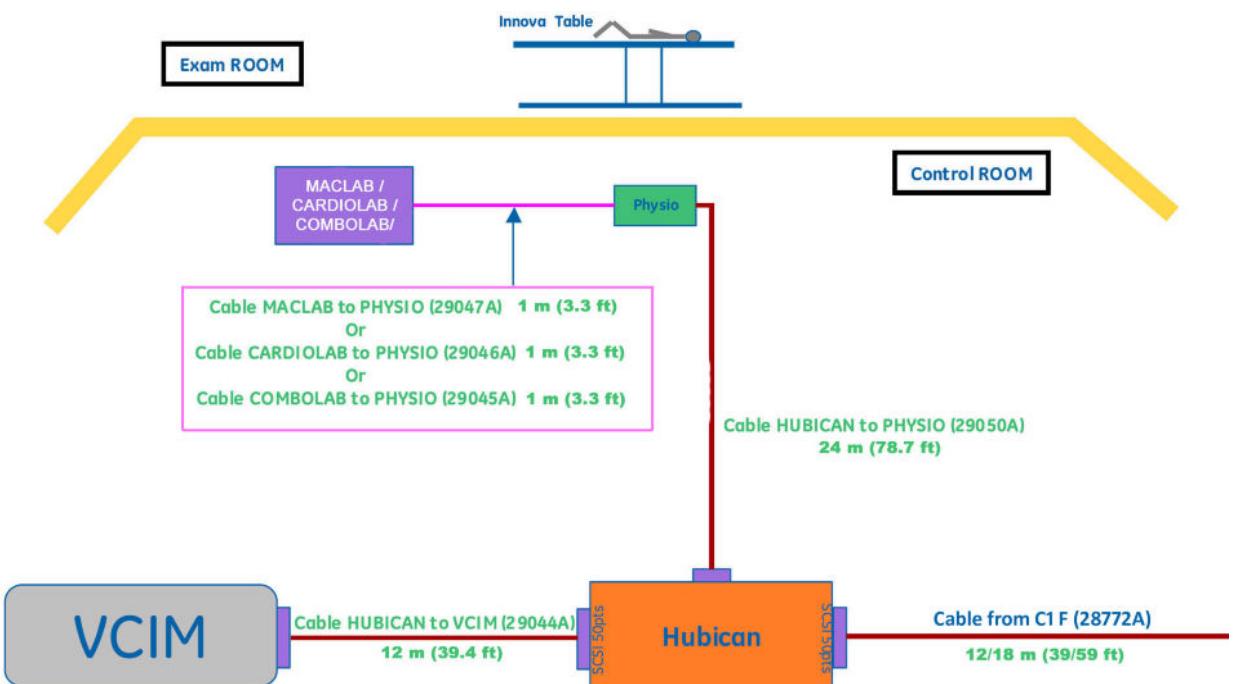
Basically, there will be two different Installation Configurations depending on the type of ECG device to connect to the Innova system:

2.2.5.1 Installation Configuration #1:

For ECG devices installed in Control Room (such as GE ECG like MacLab, CardioLab or ComboLab)

Both the Physio & Hubican modules are installed in the Control Room as shown below:

Illustration 2-61:

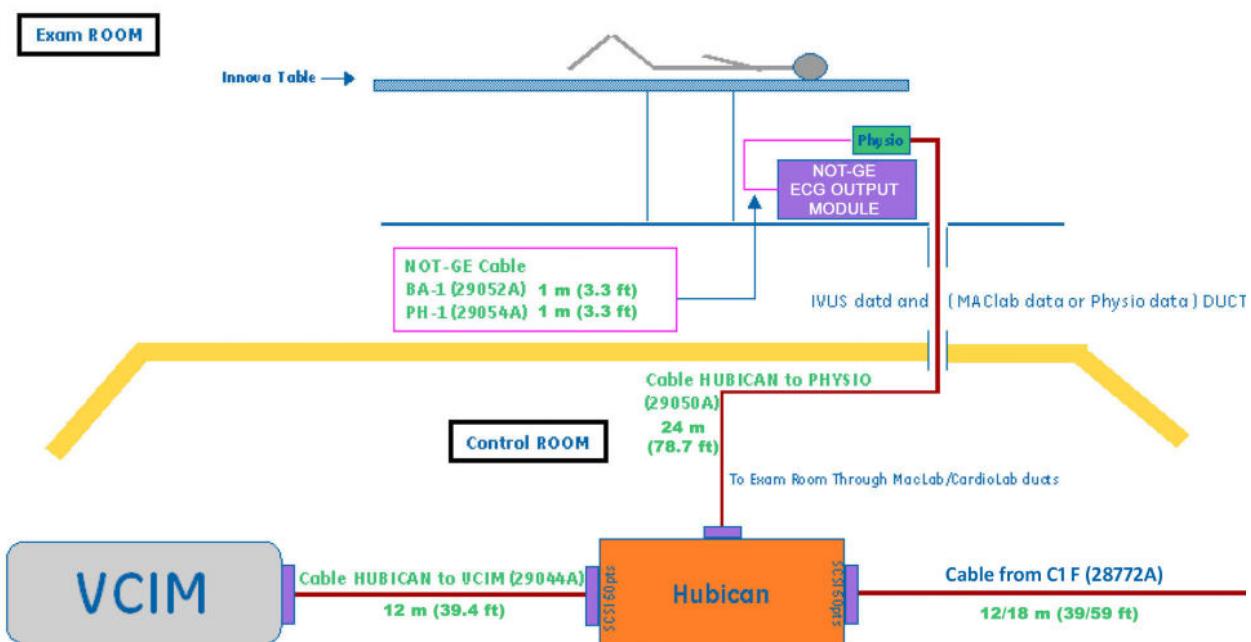


2.2.5.2 Installation Configuration #2:

For ECG devices installed in Exam Room (ECG device from the competitor)

Physio and/or Hubican modules will be installed in Exam Room as shown below:

Illustration 2-62:

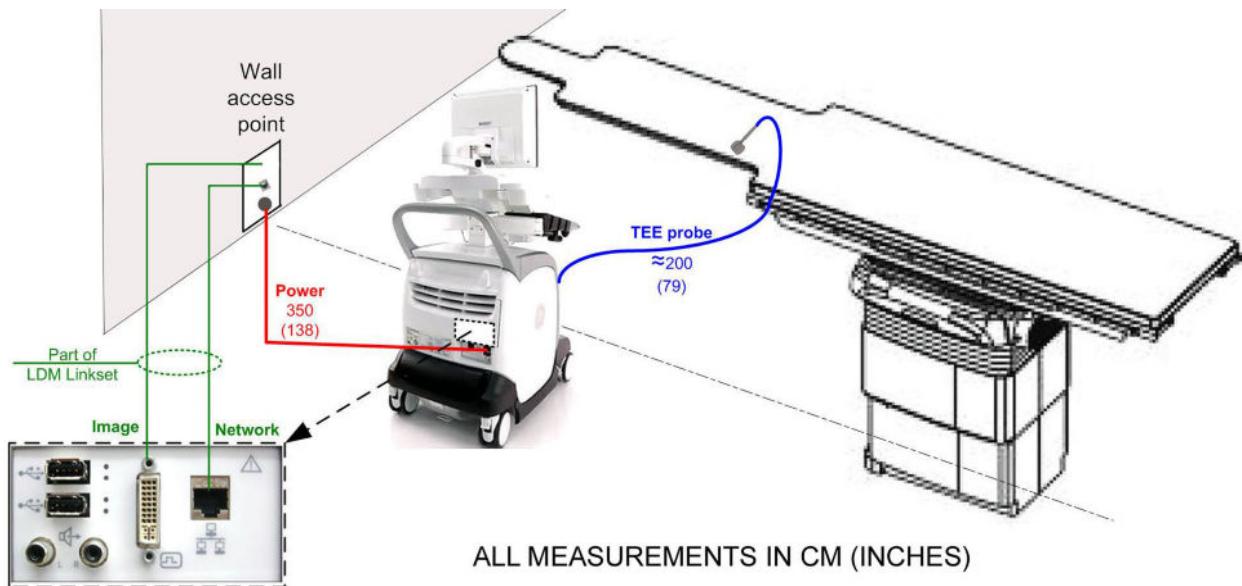


2.2.6 Vivid E9 Ultrasound Layout

On the application defined Vivid E9 positions of the Structural Heart procedures, the 3.5 m power connection cord limits the location of the power access point. For the minimal perturbation of the layout, the 3 Vivid E9 connection cables (power, network and the LDM image link) shall run close to each other, and shall be directed to a connection area on the head end wall of the patient room as close to the floor as it is possible, in order to keep the cables on the room floor.

For information concerning cable conduit from technical room to head end wall of the patient room, refer to [Chapter 5, Cable Channeling](#).

Illustration 2-63: Vivid E9 Layout (Positioner not shown)



NOTE: The power and the network connection cables belong to the Vivid E9, and they shall be disconnected from the wall access sockets before moving the scanner out of the application positions.

On the other hand, the **LDM image link cable shall be disconnected from the back of the Vivid E9**, and rolled up on the cable hooks on the wall.

2.3 Room Layout Considerations

2.3.1 Radiation Protection

Because X-Ray equipment produces radiation, you may need to take special precautions or make special site modifications. The General Electric Company does not make recommendations regarding radiation protection. It is the purchasers' responsibility to consult a radiation physicist for advise on radiation protection in x-ray rooms.

2.3.2 Service Access

Allow appropriate space for service access of equipment. Consult component pre-installation directions for clearance information.

2.3.3 Clinical Access

Make sure that you plan the room with the following clinical access requirements:

- Provide easy access to the patient table. Stretchers and other mobile hospital equipment must reach the table quickly.
- The layout of the table in the room (PIM) shall make a provision so that the clearance between the maximum table position (head side) on system axis and any object in the room (e.g.: wall, device) be greater than 50 cm (19.7 in) or 65 cm (25.6 in) if the Header Extender is used), taking into account the fact that the Tilting table can rotate 180°.
- Provide sufficient space around the patient table for the unimpeded conduct of CPR (Cardiac Pulmonary Resuscitation). With the table in this position, the table must be capable of rotating ± 45°
- Clinicians at the patient table must be able to communicate with assistants in the control area.
- There must be an unrestricted view of the video monitors and physiological monitoring equipment from the vascular table.
- Operators in the control area must have easy access to the control console. However, position the controls (including handswitches) so that the operator cannot take exposures while looking around or standing outside the control booth's lead glass window.
- Operators in the control area must have easy access to video recorders and injector programmers, film and video storage cabinets, and service and operating manuals.
- Consult customer on the number and location of nonelectrical lines (air, oxygen, vacuum, water, etc.) in the vascular room.
- For Large Display systems, in case of failure of the main monitor, the clearance around the main monitor suspension must assure that it can be immediately be flipped at 180°, exposing the backup monitors.
- For Vivid E9 Option, during the clinical procedures the Vivid E9 will be positioned either on the left, or on the right side of the patient head area, at a distance which permits comfortable access of the TEE probe towards the patient. The Vivid E9 operator stands between the equipment and the patient head. The patient head area positions are shared with the anesthesiology equipment and operator, and the interventional cardiologist and/or surgeon.

During the X-Ray operation, the Vascular gantry movements also limit the Vivid E9 positions, and a dedicated back out path has to be cleared for Vivid E9 park position.

2.3.4 Peripheral Equipment

Consult hospital personnel regarding additional space requirements for the following types of hospital equipment:

- Storage cabinets.
- Sinks.
- Oxygen stations.
- IV apparatus.
- Injectors.
- Heart monitoring equipment.
- Crash cart.

2.3.5 Emergency Stop

Protect the Emergency Stop from accidental actuation.

2.3.6 Patient Environment Equipment



WARNING

OPTIONAL LARGE DISPLAY SECONDARY MONITOR ON WALL-MOUNTED SUSPENSION MUST BE INSTALLED OUTSIDE OF THE PATIENT VICINITY

The components that may be installed within patient vicinity need to be medical equipment ("patient vicinity is defined in the standardization as a space within the room 1.83 m (70.7 in) beyond the perimeter of the examination table and extending vertically 2.29 m (90.2 in) above the floor."). For the Innova System, the equipment are:

- Table
- C-arm
- Monitors
- Injector
- Rad-Shield
- Intra Vascular Ultrasound System (IVUS)
 - IVUS Control Console.
 - IVUS Joystick.
 - IVUS Touch Pad

- Patient Interface Module(s).
- Table Side User Interfaces (TSUI)
 - ISmart Box
 - Table Side Status Control
 - Table Panning Handle
 - ITU
- NOTE: No IPX4 components shall be installed on a cart.
- Innova Central touch screen
- In-room 3D Mouse
- Accessories:
 - Table Head Extender



! WARNING

THE TABLE HEAD EXTENDER SHALL NOT BE USED WHILE THE TABLE IS TILTED. FAILURE TO DO SO MAY RESULT IN SEVERE PATIENT INJURY.



! WARNING

THE TABLE HEAD EXTENDER IS NOT COMPATIBLE WITH THE INNOVA IQ WIDE TABLE TOP.

- Armboard
- IV pole
- Quick Strap
- Head holder: Attach Head Positioning device onto the head end of the table with the head cradle pointing to the end of the table. The clamp angle pressure bar should be resting on top of the table and the blue stabilizing feet under the table. Tighten the clamping systems with the four- pronged knob. Test to check that head holder is securely attached to the table. Adjust the height and angle of the head cradle by loosening the long handled knob. Once cradle is set at desired position tightened the long handled top. Place the patient's head in the head cradle. Use the two velcro© straps to hold the forehead and chin in place. Additional adjustment maybe made to the position of the head cradle by loosening and tightening the long handled knob if desired.



⚠️ WARNING

THE HEAD HOLDER SHALL NOT BE USED TO RESTRAIN THE PATIENT'S HEAD WHEN THE TABLE IS TILTED, ONLY WHEN IT IS HORIZONTAL. IT IS NOT DESIGNED TO SUPPORT THE HEAD FROM MOVING AND MAY CAUSE INJURY TO THE PATIENT.

- Clear-Vu Arm Support
- Mattress and mattress slicker
- Shoulder Rests
- Footrest
- Removable rails (sleeve)



⚠️ WARNING

SLEEVE IS NOT COMPATIBLE WITH INNOVA IQ WIDE TABLETOP & WITH HEAD EXTENDER.

- Rail Extender
- Head Widener



⚠️ WARNING

HEAD WIDENER IS NOT COMPATIBLE WITH HEAD EXTENDER & SHOULDER REST.

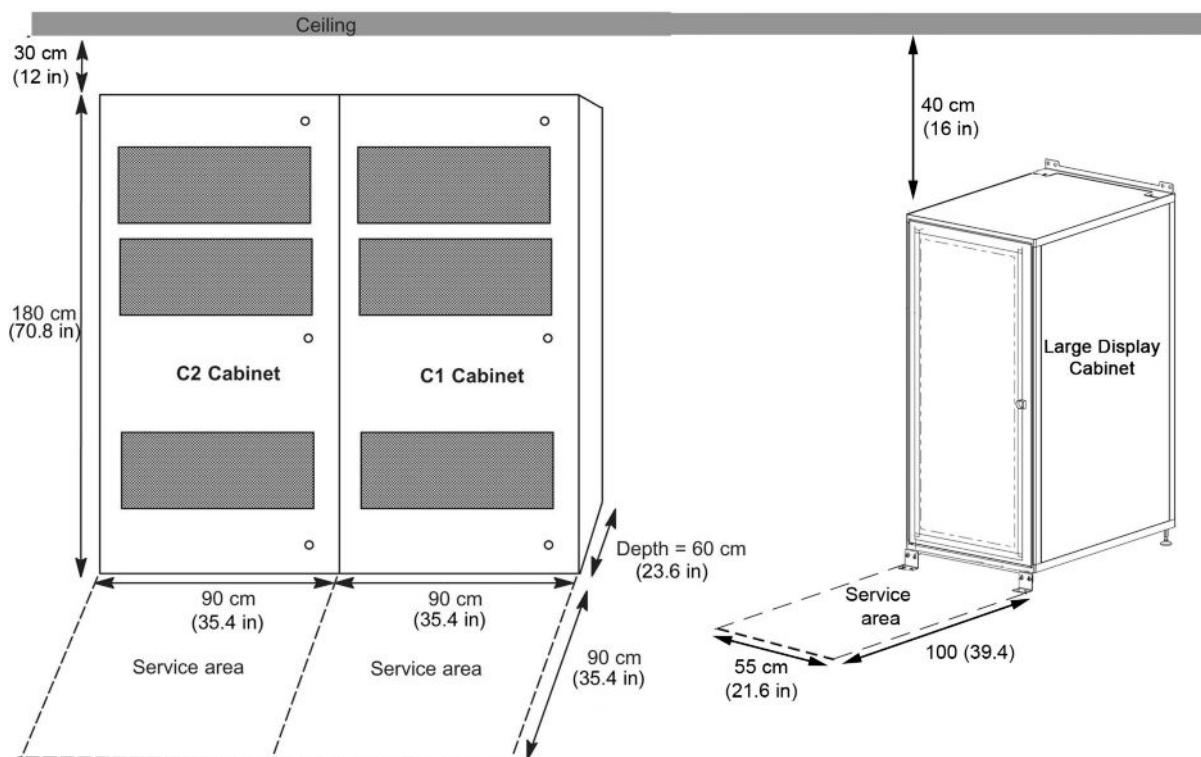
- Adaptor rail for table side controls
- Width Extender
- Armboard with thick pad (armrest)
- Universal Clamp
- Patient restraint strap

2.3.7 Preference Cabinet locations

NOTE: This is applicable for all types of cabinets (including PDB).

The clear width of the service area in front of the cabinet doors to insure electrical safety shall be at least 0.9 m for all cabinets except the Large Display cabinet (1 meter is necessary for Large Display cabinet). In cases where 2 cabinets are installed face to face (both sides of the access way), the clear width shall be at least 1.2 m.

Illustration 2-64:



CAUTION

The service area dimensions shown above are minimum requirements. Service areas must comply with local regulations if more stringent.

2.3.8 Layout Constraints



NOTICE

The X-Ray tube Chiller elevation cannot exceed 3 meters (9.84 feet) in height.



NOTICE

Minimum distance between Digital Detector and UPS is 3 m (118 inches) (Fluoro UPS option).



NOTICE

The distance of the Detector chiller cannot exceed 3 m (1 floor) in height below the detector.

3 Room Structural Requirements

3.1 General Policy

GE Healthcare's Customer is responsible for the structural analysis and mounting of the base plates. If GE Healthcare is forced to mount the base plate, the LCT must hire a structural engineer to design and approve the mounting method and provide GE Healthcare with an engineering report.

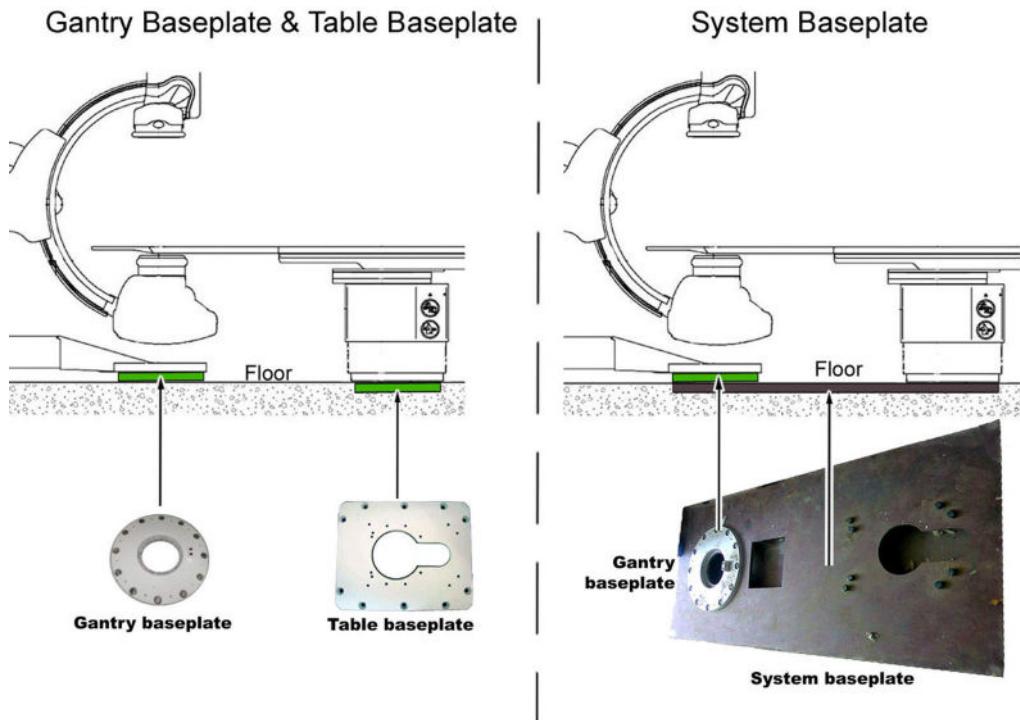
The floor level cannot exceed a general levelness of 5 mm (0.2 in) for any 2 meters (79 in).



NOTICE

The floor slabs on which the equipment is to be installed must have a levelness of 1 mm (0.04 in) per meter (40 in). Position of baseplates and table basement depends on the type of installation. The two types of installation are given below.

Illustration 2-65: Types of baseplate installation



NOTICE

The Table baseplate or the System baseplate are mandatory to install the Tilting table.

The Tilting table must never be installed on grade.



NOTICE

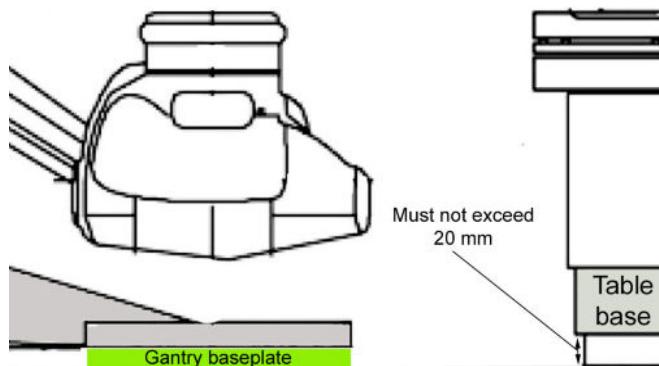
Augmented Calibration requires that the patient table and gantry be installed following Pre-installation instructions. Any deviation may result in a non-augmented calibration.



NOTICE

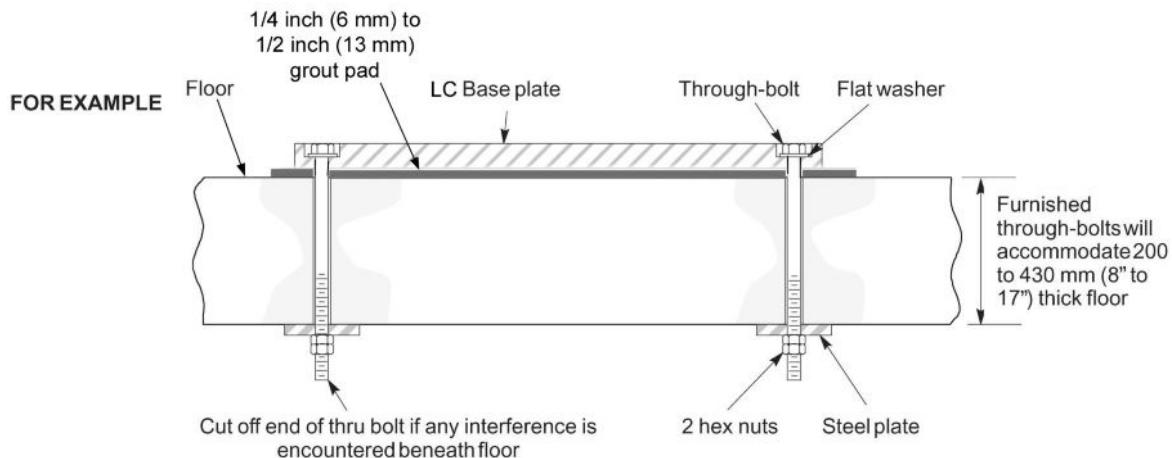
The gap between the Table Foot bottom end the Gantry Baseplate bottom end shall be lower than 20 mm (0.97 in). Any bigger gap would make the system incompatible with the Innova Vision Applications.

Illustration 2-66: Gap between Table Foot bottom end the Gantry Baseplate bottom



The preferred installation method for the Innova LC Positioner or the Omega tables is through-bolting. The through-bolting method can be used in all seismic zones. If through-bolting cannot be used, use provided floor anchors instead.

Illustration 2-67: Through-Bolt Supplied (Slab Type Floor Construction)



3.2 Floor Requirements

3.2.1 Floor requirements when using provided floor anchors

The maximum pullout force per provided anchor was calculated assuming:

- A concrete compression strength of **17.24 MPa** at 28 days (which is the minimum required compression strength).
- Anchors installed to the required hole depth of **165.1 mm** minimum, and
- Center of anchor hole to concrete edge distance **79.4 mm**.

Make sure to obtain data on compression strength of the concrete before using floor anchors.

3.2.2 Pan Type Floor Construction Requirement

For Pan type floor construction, steel channels must be designed by a local structural engineer to span floor joists. See [Illustration 2-68](#).

NOTE: For specific floor preparation procedures, refer to *Pre-Installation Kit Installation Procedures*.

Illustration 2-68: Through-Bolt Supplied (Pan Type Floor Construction)

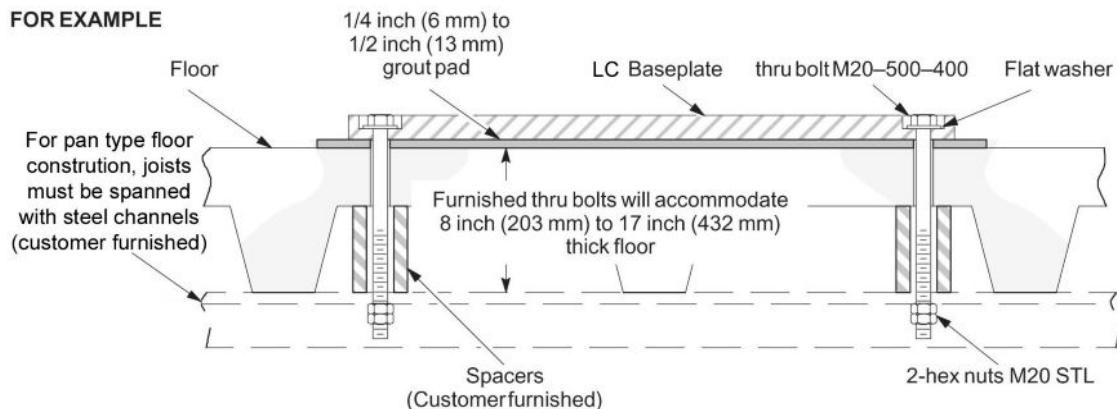
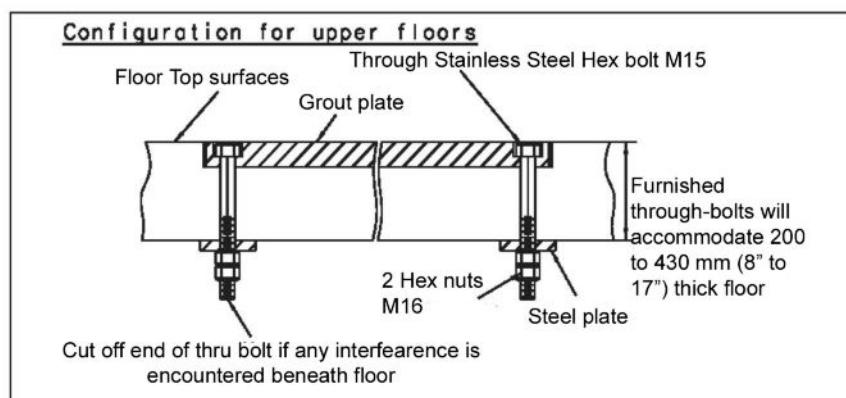
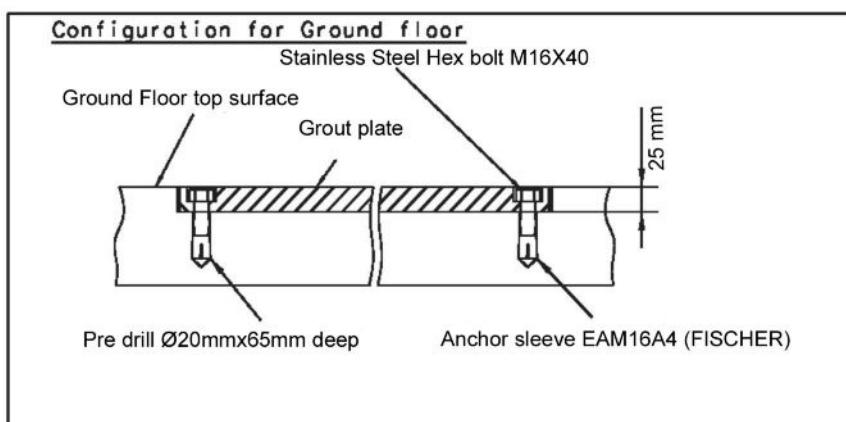


Illustration 2-69: Table floor mounting layout



NOTICE

Prepare the floor such that the table base plate will be flush with the floor finish surface, taking into account the thickness of the floor finish material.

For alternative table bolts or seismic area, refer to template drawing shown in Illustration *Innova LC Positioner And Table Floor Mounting Template* contained in [Mounting Requirements](#).

3.2.3 Hole dimension and preferred location in concrete floor

In the examination room, the Innova LC Positioner is not placed on a computer floor but directly put on concrete floor, the location of the cable access needs to be carefully planned.

Otherwise, if the cable run is located under the concrete floor, the cables will have to come through the floor and in this case you will need two holes, one for the LC Positioner and the other for the patient table.

The diameter of both holes must be the same 225 mm or 9 in.

Illustration 2-70: Hole location in concrete floor

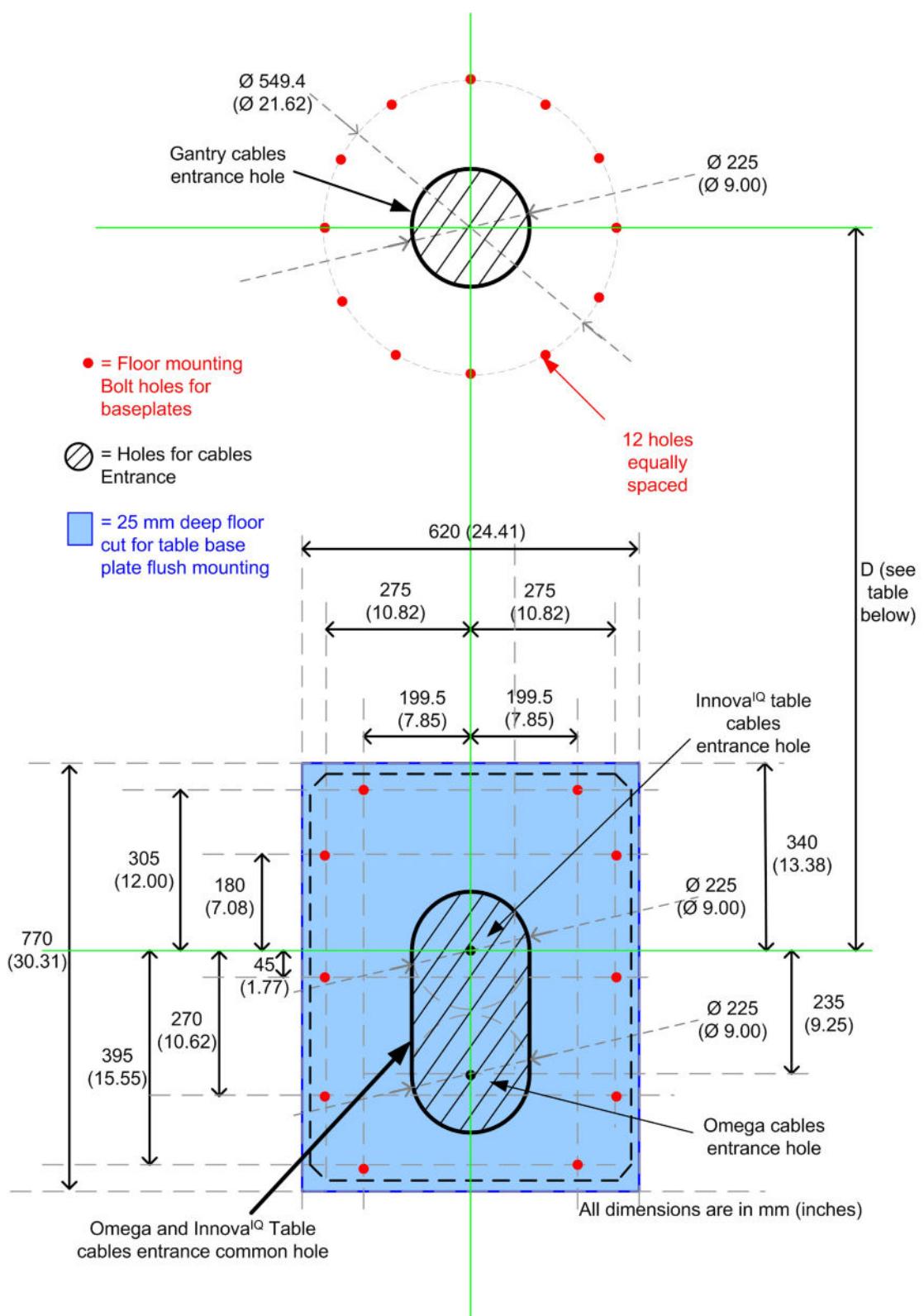


Table 2-7: D distance

	ANGIO	CARDIO	NEURO
Omega IV Compact	NA	1395 mm (54.9 in)	NA
Omega V Long	1278 mm (50.3 in)	1395 mm (54.9 in)	1395 mm (54.9 in)
Omega V non motorized Long	1278 mm (50.3 in)	1395 mm (54.9 in)	1395 mm (54.9 in)
Tilting table	1278 mm (50.3 in)	1395 mm (54.9 in)	1595 mm (62.8 in)



NOTICE

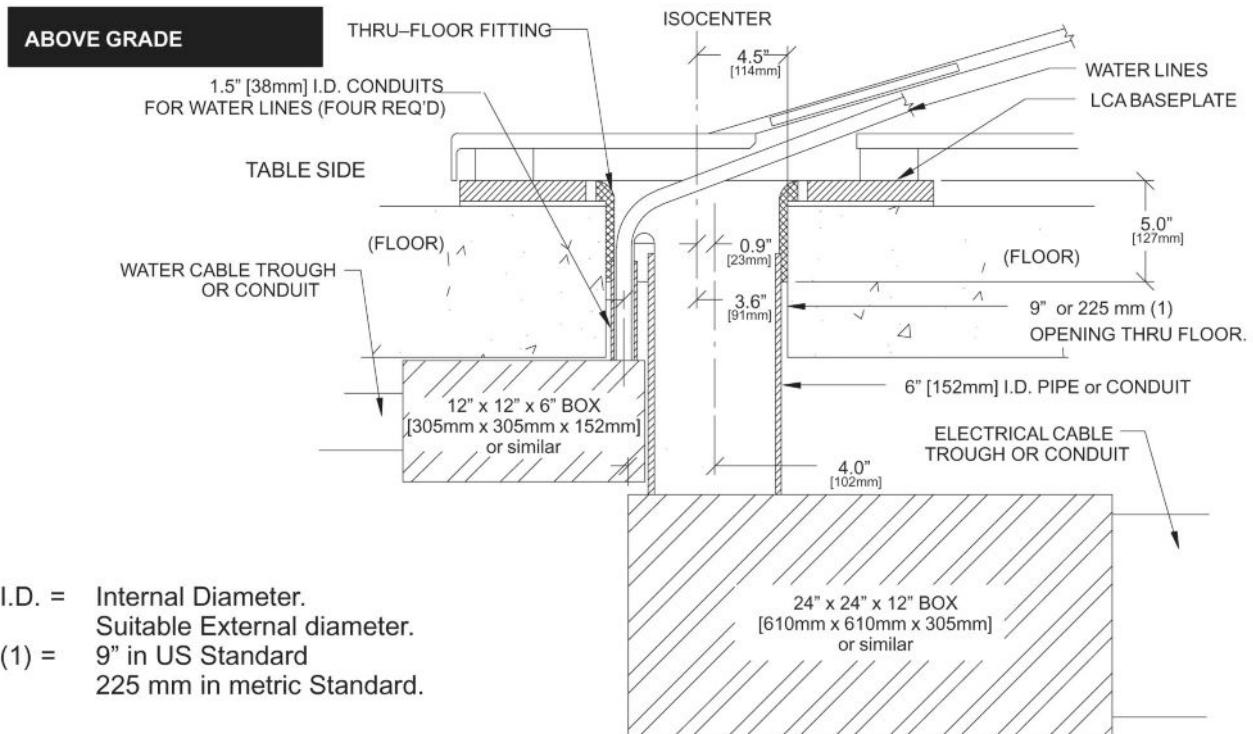
Due to the plastic bushing used in the USA to protect cables from the sharp edges of conduits it is necessary to place the cable conduit inside the table cable access opening but the height of the outgoing conduit plus bushing is limited to 1/2 in (12.7 mm).

NOTE: Refer to table *Chemical anchors Pull out efforts and recommendations* in [Mounting Requirements](#) for pull out effort on each fixation bolts.

3.2.4 Water Pipe Requirements

- An Innova LC System uses a Performix 160 A X-Ray tube with a recirculating chiller.
- Two water hoses are supplied to allow water circulation between Innova LC Positioner and chiller.
- In some countries, it is forbidden to run electrical cables and water pipes in the same conduit. In this case, two separate conduits are required. But then a problem arises at the level of the Innova LC Positioner entrance.
- Depending on the method used, one of the two options shown hereafter ([Illustration 2-71](#) and [Illustration 2-72](#)) must be used.

Illustration 2-71: Water Conduit location with “Above Grade” anchor kits



I.D. = Internal Diameter.
 Suitable External diameter.
 (1) = 9" in US Standard
 225 mm in metric Standard.

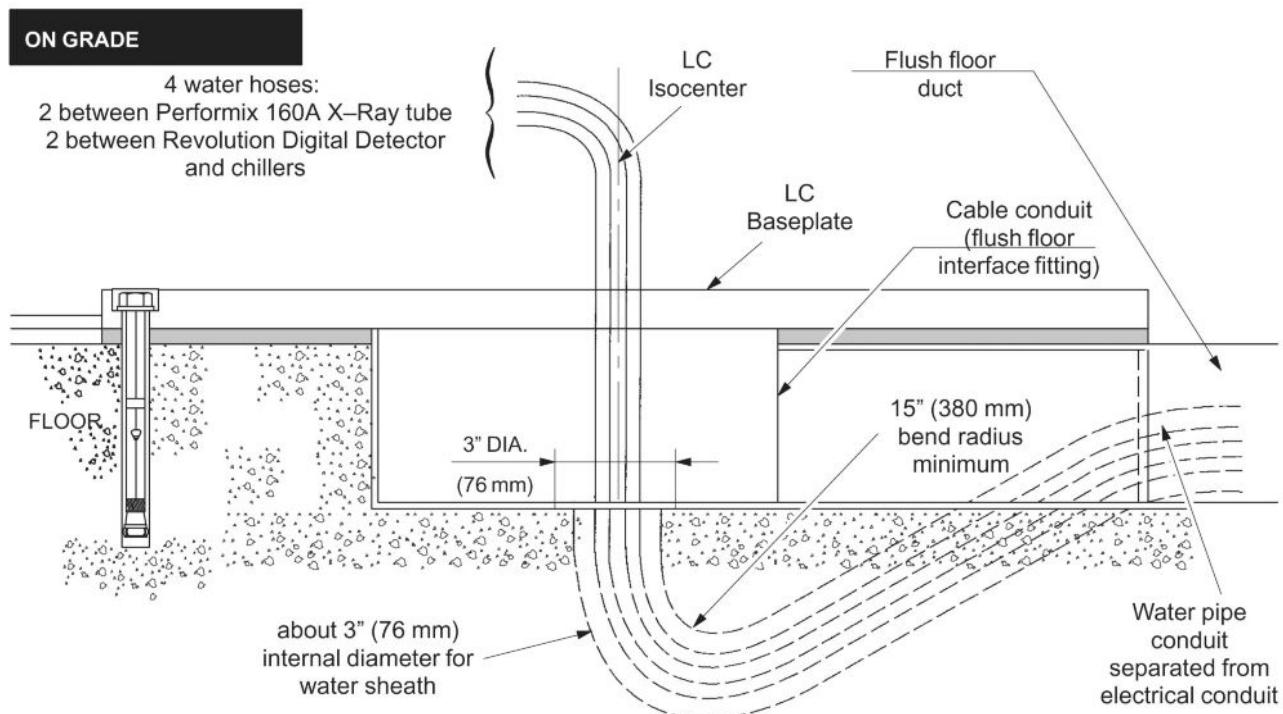
Note: Pipe, junction box and duct or conduit are to be supplied and installed by Customer or customer's Contractor.



NOTICE

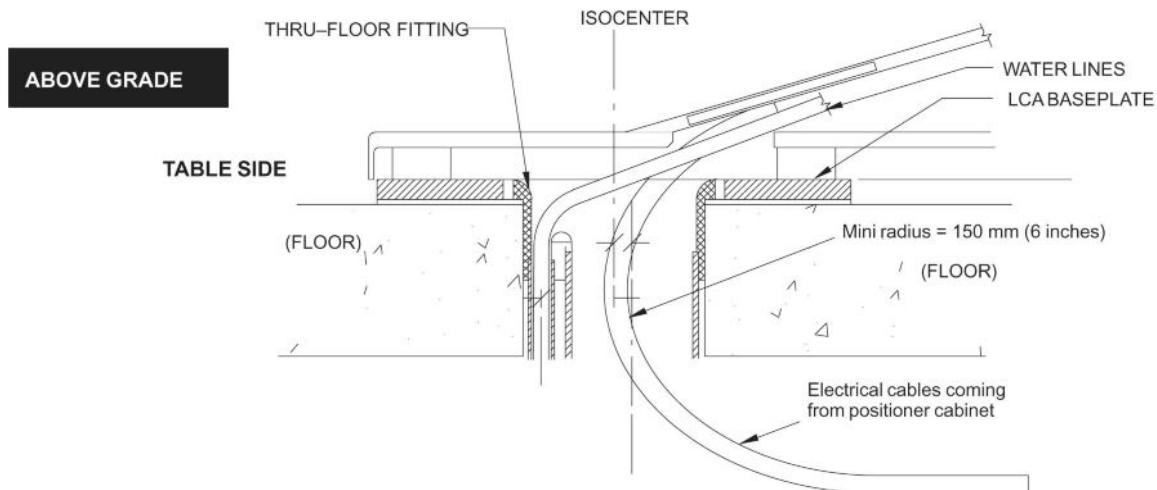
In some countries, depending on local regulations, it may be forbidden to run electrical cables and water pipes in the same conduit. In this case, two separate conduits are required.

Illustration 2-72: Water Conduit location with “On Grade” anchor kits



Note: Flush floor interface fitting is part of GEMS installation kit 2286398 and is installed by Customer or customer's Contractor.

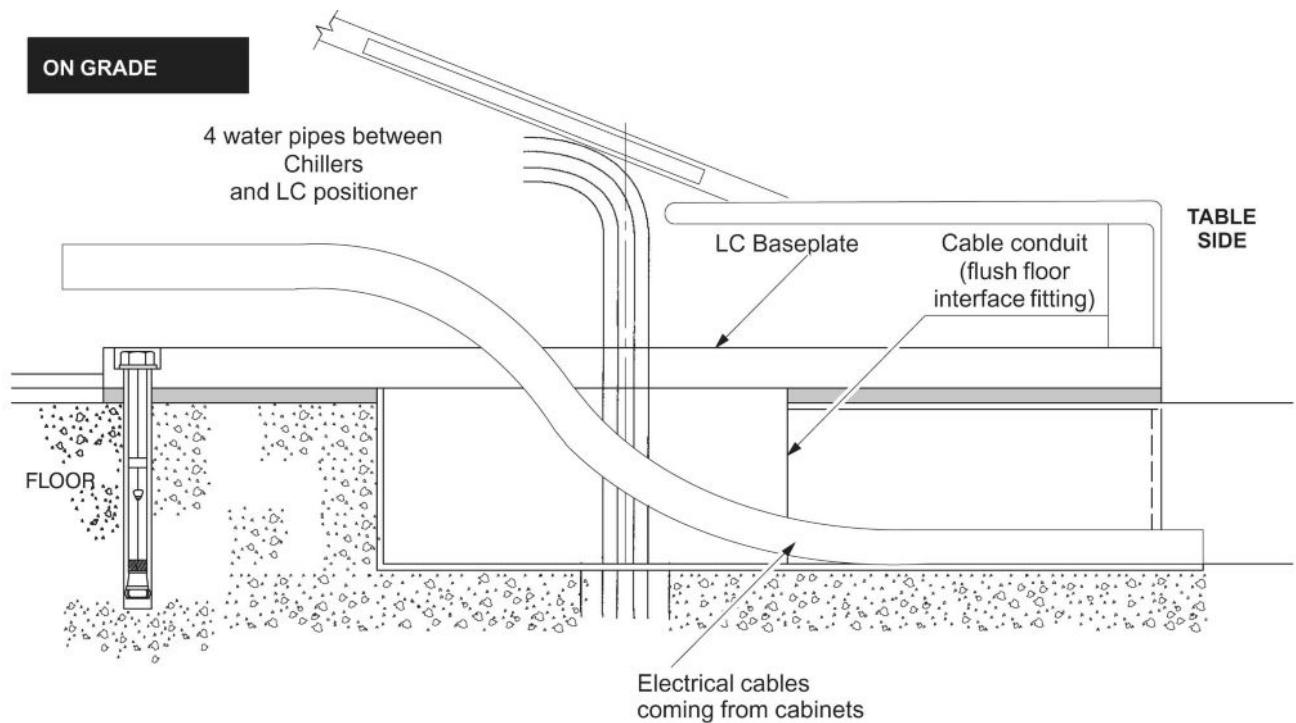
Illustration 2-73: Cable Curvature with “Above Grade” anchor kits



Note: In case of thru-floor cabling, if the electrical cables are coming from the head side, they will need to have a minimum curvature with a minimum radius of 150 mm (6").

In any other cases (i.e. flush floor) no such curvature is allowed.

Illustration 2-74: Cable Curvature with “On Grade” anchor kits



NOTE: In case of on grade cabling, because of the minimum curvature constraint of 150 mm (6"), the cable will have to come from the side between Innova LC Positioner and patient table.

3.3 Mounting Requirements

3.3.1 Floor Loading and Recommended Mounting Methods

See [Table 2-8](#). To obtain floor loading and recommended mounting methods for components not specified in [Table 2-8](#), refer to the appropriate component Pre-Installation Manual listed in [Basic Innova System Compatibility](#).

Table 2-8:

PRODUCT OR COMPONENT	NET WEIGHT KG (LBS)	DIMENSIONS MM (INCHES)			LOAD BEARING AREA MM (INCHES)	WEIGHT/ OCCUPIED AREA	MOUNTING METHOD
		WIDTH	DEPTH	HEIGHT			
Gantry	750 (1655)	See Illustrations <i>GANTRY DIMENSIONS</i> : • Side view • Top view • Front view in Dimension Drawings	Circle diameter 600 (23.62)			Recommended: <ul style="list-style-type: none">Through-Bolts (12) Alternates: <ul style="list-style-type: none">On Grade 5/8 in, Anchors (12)Above Grade 3/4 in. Anchors (12) See Illustration 2-75 , Illustration 2-76 and Illustration 2-77	
Table	590 (1300) See NOTE (1)	See illustrations in Dimension Drawings • Omega IV • Omega V	571.5x429 (22.5x16.9)	2410 kg/m ² (492.3 lb/ft ²)	Same as Gantry See IMPORTANT NOTICE below.		
	785 (1731) See NOTE (1)	Tilting table	750x600 (30x24)	2000 kg/m ² (409 lb/ft ²)			
C2 Cabinet	258.5 (570)	See Illustration <i>C2 Cabinet Dimensions</i> in Dimension Drawings	600x900 (23.63x35.44)	478 kg/m ² (98 lb/ft ²)			
C1 Cabinet	457 (1007.5)	See Illustration <i>C1 Cabinet Dimensions</i> in Dimension Drawings	600x900 (23.63x35.44)	838 kg/m ² (172 lb/ft ²)			
COOLIX 4100	120 (264.5)	555 (21.8) 610 (24) 1200 (47.2)					
Autotransformer (Coolix 4100)	30 (66)	370 (14.5) 304 (12) 340 (13.4)					
Detector Chiller Thermo-Con	14.6 (32.2)	See Illustration <i>Detector Chiller Thermo-Con Dimensions</i> in Dimension Drawings	344x283 (13.5x11.14)				
Fluoro UPS UL (optional)	530 (1169)	690 (27.0) 800 (31.5) 1820 (70.1)			975 kg/m ² (200 lb/ft ²)	Ground -mounted	
Fluoro UPS CE (optional)	480 (1059)	680 (26.7) 800 (31.5) 1450 (57.1)			883 kg/m ² (181 lb/ft ²)	Ground -mounted	
Fluoro UPS IF box (optional)	4 (9)	280 (11.02) 314 (12.36) 124.5 (4.9)				Wall-mounted	
3 kVA UPS	34.5 (76.1)	See <i>3 kVA UPS Dimensions</i> in Dimension Drawings				Ground -mounted	

PRODUCT OR COMPONENT	NET WEIGHT KG (LBS)	DIMENSIONS MM (INCHES)			LOAD BEARING AREA MM (INCHES)	WEIGHT/ OCCUPIED AREA	MOUNTING METHOD
		WIDTH	DEPTH	HEIGHT			
PDB US (4)	148 (326)	685 (27)	225 (9)	1850 (73)			Wall-mounted
PDB CE (4)	110 (242)	800	300	1200			Wall-mounted
EMI Filter enclosure CE (4)	<40	300	210	950			Wall-mounted
DL keypad	1.4 (3)	283 (11.55)	300 (11.8)	82 (3.25)			
DL LCD monitor	8.2 (18)	179 (7)	387 (15.2)	504 (19.8)			
VCIM	0.95 (2.09)	450 (17.7)	150 (5.9)	50 (2)			
Videostation VCR	12 (26.5)	370 (14.57)	270 (10.63)	144 (5.67)			
Large Display monitor suspension (See NOTE 5)	356 (784)	See illustration <i>Large Display suspension dimensions (Optional)</i> in Dimension Drawings					Ceiling-mounted
IVUS S5I GE Rev 2 Option							
PIM (Ultrasound probe)	2 (4.4)	38	96	177			
Control Station	4 (8.8)	120	360	240			
Joystick	1 (2.2)	90	76	102			
Isolation transformer	7 (15.4)	97	267	171			
IVUS CPU	15 (33.1)	396	162	510			
IVUS Keyb. & mouse pad	1 (2.2)	NA	NA	NA			
IVUS monitor	15 (33.1)	420	432	240			
Image printer	6 (13.2)	158	184	18			
4-way Video Switch	1 (2.2)	44	222	241			
PIMr	1,5 (3,2)	356	102	102			
Pimette	1 (2,2)	142	76	18			
Touchpad controller	1 (2,2)	211	180	76			
Printer HP	1,3 (2,8)	227	138	117			
LD cabinet	115 (254)	550 (22)	780 (31)	1250 (49)	550x780 (22x31)	268 kg/m ²	
Vivid E9 (See NOTE 6)	128 (283)	540 (21.2)	800 (31.4)	1375/1575 (54.1/62.0)	Not Applicable	Not Applicable	Not Applicable
IVUS S5I GE Rev 2+ Option							
PIM (Ultrasound probe)	0.7 (1.5)	91 (3.58)	212 (8.35)	41 (1.61)	41 x 212 (1.61 x 8.35)	N/A	In holder
PIMr	1.5 (3.3)	102 (4.02)	366 (14.41)	99 (3.90)	102 x 366 (4.02 x 14.41)	0.4 kg/m ²	Placing on table
Pimette	0.5 (1.1)	77 (3.03)	144 (5.67)	36 (1.42)	77 x 144 (3.03 x 5.67)	0.45 kg/m ²	On IV pole

PRODUCT OR COMPONENT	NET WEIGHT KG (LBS)	DIMENSIONS MM (INCHES)			LOAD BEARING AREA MM (INCHES)	WEIGHT/ OCCUPIED AREA	MOUNTING METHOD
		WIDTH	DEPTH	HEIGHT			
Control Station	3.6 (7.9)	330 (12.99)	177 (6.97)	100 (3.94)	330 x 177 (12.9 x 6.97)	0.6 kg/m ²	On desk or rail
Joystick	0.5 (1.1)	90 (3.54)	76 (2.99)	102 (4.02)	N/A	N/A	On rail
Touchpad controller	1 (2.2)	211 (8.31)	180 (7.09)	76 (2.99)	N/A	N/A	On rail
Isolation transformer	6 (13.2)	165 (6.50)	267 (10.51)	89 (3.50)	165 x 267 (6.50 x 10.51)	1.4 kg/m ²	On floor
S5I GE CPU	18 (39.7)	522 (20.55)	176 (6.93)	430 (16.93)	176 x 522 (6.93 x 20.55)	2.0 kg/m ²	On floor
S5I GE Keyb. & mouse pad	1 (2.2)	N/A	N/A	N/A	N/A	N/A	On desk
S5I GE monitor	9 (19.8)	330 (12.99)	247 (9.72)	596 (23.46)	330 x 247 (12.99 x 9.72)	1.1 kg/m ²	On desk
Image printer	6 (13.2)	158 (6.22)	184 (7.24)	180 (7.09)	158 x 184 (6.22 x 7.24)	2.1 kg/m ²	On desk
Printer HP	1.3 (2.9)	227 (8.94)	138 (5.43)	117 (4.61)	227 x 138 (8.94 x 5.43)	0.4 kg/m ²	On desk
4-way Video Switch	1 (2.2)	220 (8.66)	241 (9.49)	44 (1.73)	220 x 241 (8.66 x 9.49)	0.2 kg/m ²	On desk
IVUS Rev 3 Option							
Refer to Volcano s5i Imaging System Integration in IGS Systems - Service Manual							

NOTE: (1) including patient weight (patient weight considered is 204 Kgs/450 lbs).

NOTE: (2) Depth.

NOTE: (3) Maximum dimensions given. Exact dimensions depend on chiller manufacturer type.

NOTE: (4) Recommended supplier.

NOTE: (5) The total weight (Large Display monitor suspension boom (spring arm), frame, precabled, bridge, rails monitors (LDM + 2 B/W) and accessories) is calculated as follows:

110 kg (frame+arm+monitors max load) + 50 kg (internal cabling) + 31 kg (bridge plate + main bearing + connection box) + 102 kg (XT long bridge) + 63 kg (2 stationary rails 19") = 356 kg (784 lbs)

NOTE: (6) Vivid E9 weight and dimensions include monitor and peripherals.

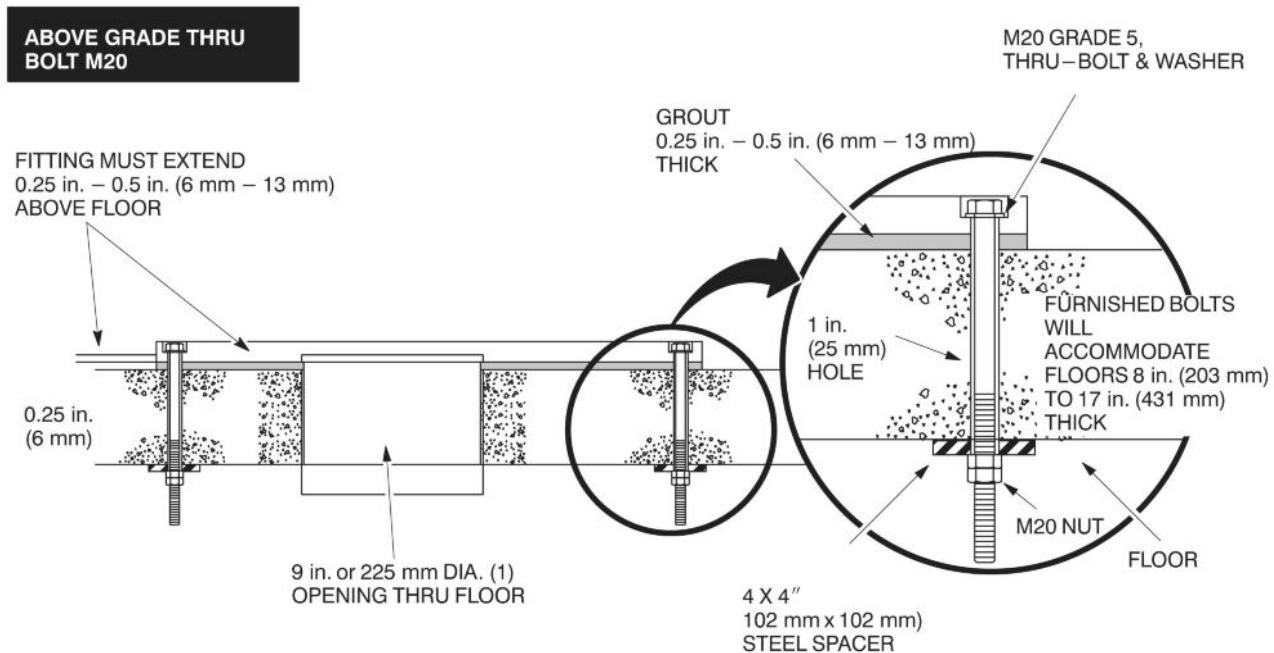
3.3.2 Positioner and Table Floor Mounting

The distances between the Gantry and the Tables are critical for a proper clinical usage. For this reason, GEMS provides two floor mounting templates to ensure these components are properly placed in relation to one another.

Table 2-9:

Title	Illustration
Gantry Floor Mounting Methods	Illustration 2-75 and Illustration 2-76
Inner Base Plate (Water Electric Separator) For Above Grade Floor Anchor Kit	Illustration 2-77
Cable Conduit For On-Grade Floor Anchor Kit	Illustration 2-78
Fixing Bolt Overview	Illustration 2-79
Gantry And Tilting table Floor Mounting Template	Illustration 2-80 and <i>Hole location in concrete floor illustration in Floor Requirements.</i>

Illustration 2-75: Gantry Floor Mounting Methods (1/2)



(1) The US or the METRIC standard for base plate inner

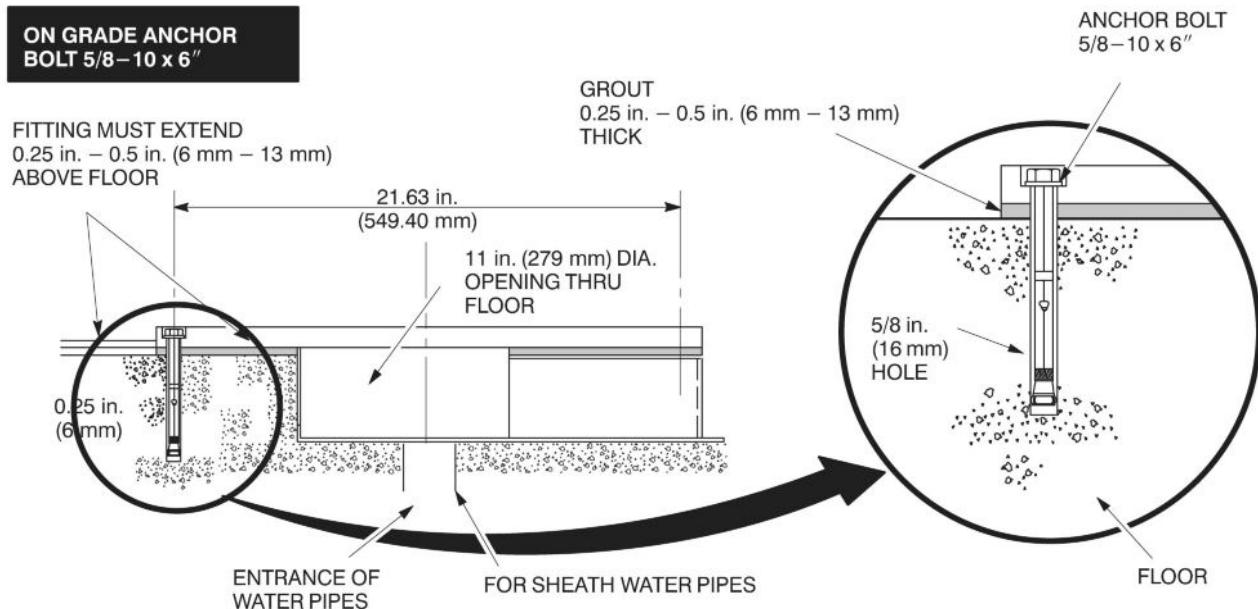
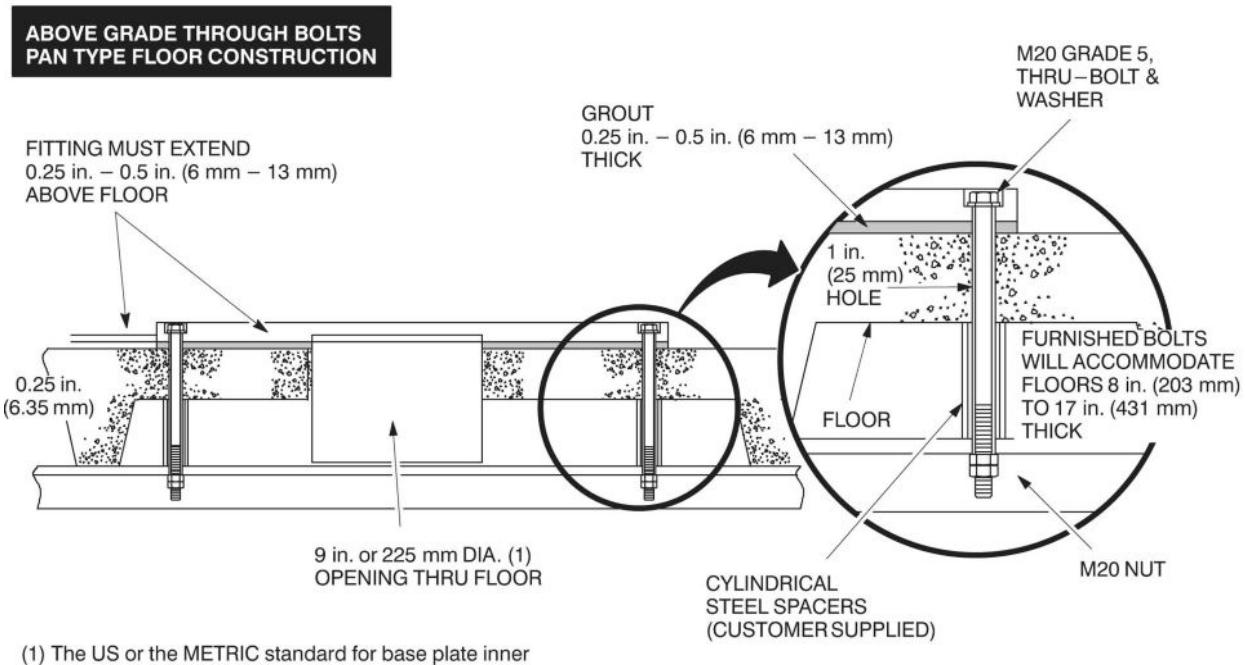
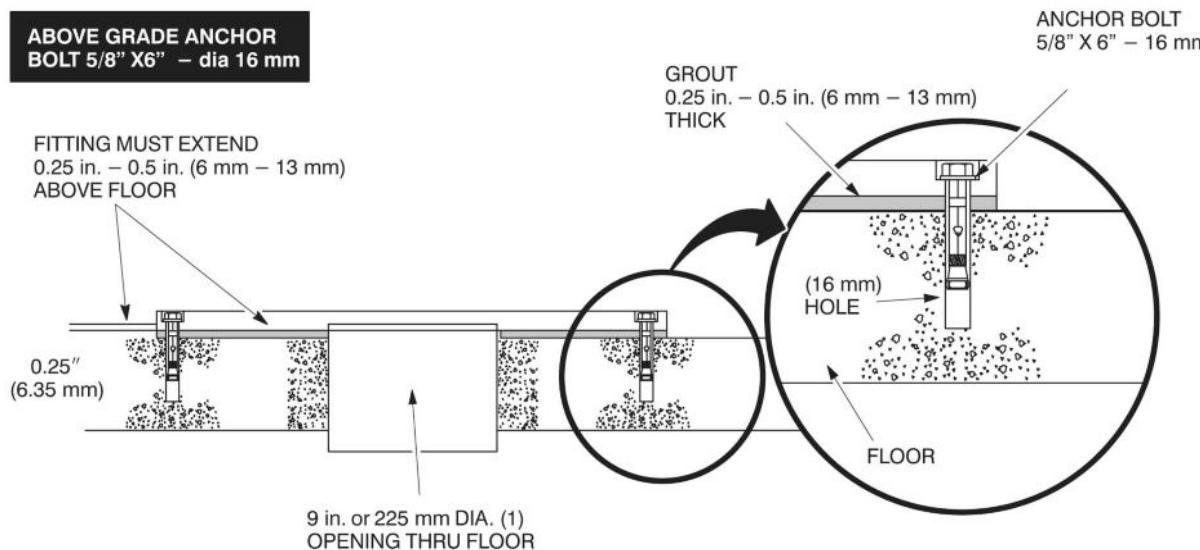


Illustration 2-76: Gantry Floor Mounting Methods (2/2)



(1) The US or the METRIC standard for base plate inner



(1) The US or the METRIC standard for base plate inner

Illustration 2-77: Inner Base Plate (Water Electric Separator) For Above Grade Floor Anchor Kit

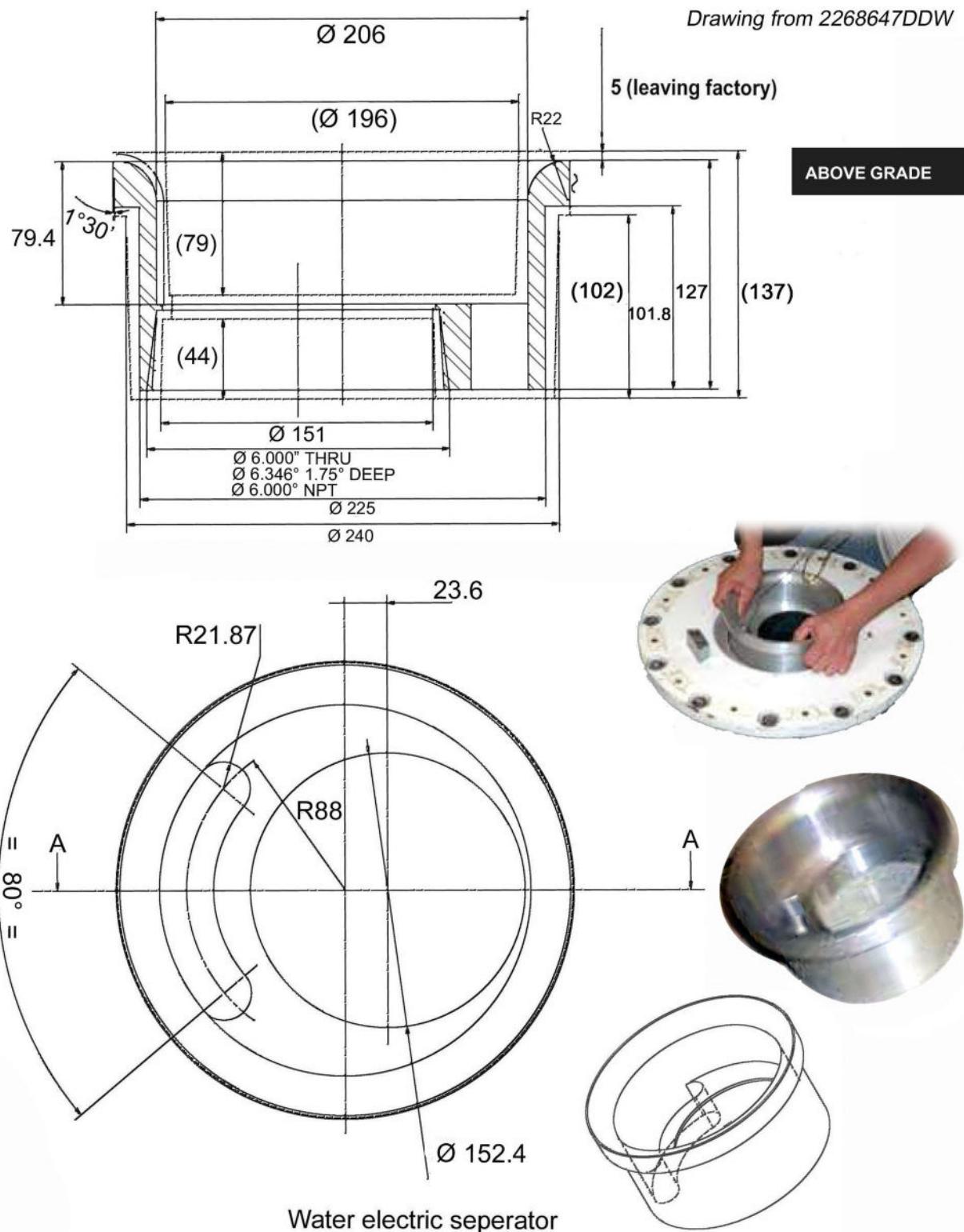
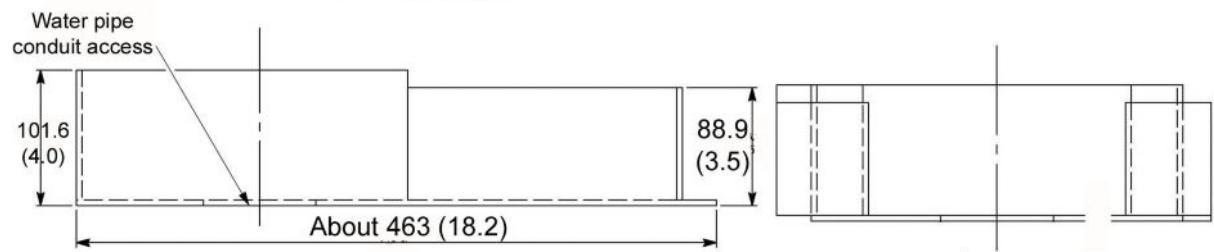
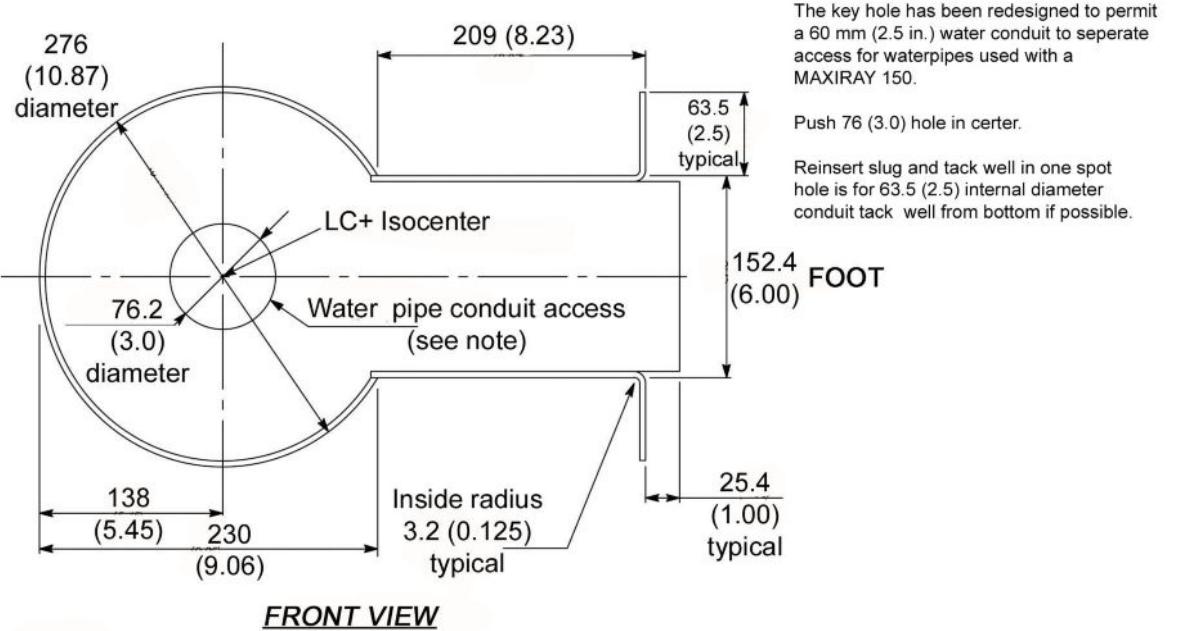


Illustration 2-78: Cable Conduit For On-Grade Floor Anchor Kit



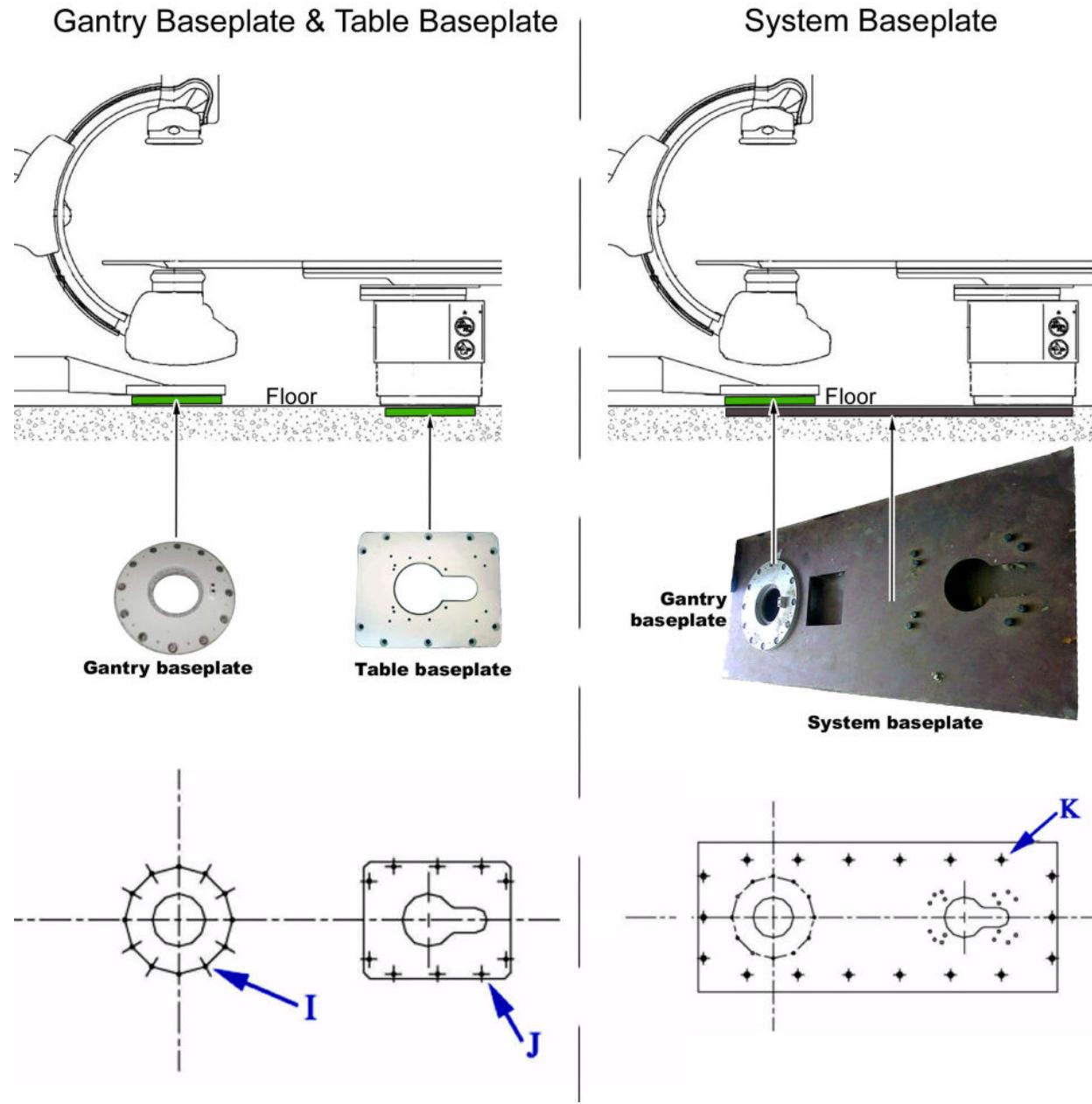
Dimensions in mm (inches)

ON GRADE



FRONT VIEW (FOOT END)

Illustration 2-79: Fixing Bolt Overview



NOTE: For more details on Floor plate or Base plates, refer to [Illustration 2-80](#) and illustration [Hole location in concrete floor](#) in [Floor Requirements](#).

NOTE: With any kind of fixation methods (Bolts M20, Mechanical anchors or Chemical anchors), the number of holes used mandatory is:

- Gantry base plate : 12 max and 8 min holes used are acceptable
- Table base plate : 10 max and 8 min holes used are acceptable
- Floor base plate : 24 max and 12 min holes used are acceptable

we can have only 2 consecutive holes omitted.

NOTE: In case of floor plate configuration, the gantry base plate shall be mounted onto the floor plate, but the table base plate shall not be mounted on the floor plate.

Pull out efforts and recommendations about chemical anchors not provided by GE.

The following table provides the recommended chemical anchors for Table/Gantry base plates and for the floor plate ordered locally that they could be used instead of bolts provided by GE.

Table 2-10: Chemical anchors Pull out efforts and recommendations

	Gantry base plate	Table base plate	Floor plate (to be ordered locally)	Table Omega
Mark	I on Illustration 2-79	J on Illustration 2-79	K on Illustration 2-79	A on Illustration 2-80
Pull out effort	736 daN per bolt if 12 used and 1992 daN per bolt if 8 used	1120 daN per bolt if 10 used and 2000 daN per bolt if 8 used	272 daN per bolt if 24 used and 2008 daN per bolt if 12 used	4432 daN per bolt with 4 bolts
Number of holes in the plate	12 max (8 min mandatory)	10 max (8 min mandatory)	24 max (12 min mandatory)	4 mandatory
Recommended chemical anchors example 1	Supplier HILTIHVU adhesive capsule + HAS Anchor rod	Supplier HILTIHVU adhesive capsule + HAS Anchor rod	Supplier HILTIHVU adhesive capsule + HAS Anchor rod	Supplier HILTIHVU adhesive capsule + HAS Anchor rod
Threaded rod	M16 A4-70 / 333 131 5/8	M20 A4-70 / 333 135 3/4	M16 A4-70 / 333 131 5/8	M20 A4-70 / 333 135 3/4
Hole diameter in the floor	18 mm (11/16 in)	24 mm (7/8 in)	18 mm (11/16 in)	24 mm (7/8 in)
Hole depth in the floor	125 mm (5 in)	170 mm (6-5/8 in)	125 mm (5 in)	170 mm (6-5/8 in)
Minimum floor thickness	180 mm (7 in)	220 mm (8-1/2 in)	180 mm (7 in)	220 mm (8-1/2 in)
Max Tightening Torque	80 N.m (59 ft-lb)	150 N.m (110 ft-lb)	80 N.m (59 ft-lb)	150 N.m (110 ft-lb)

NOTE: The floor plate ordered locally needs to be in steel.

Refer to supplier technical documents for all specification and installation data about chemical anchors.

Illustration 2-80: Gantry and table mounting holes

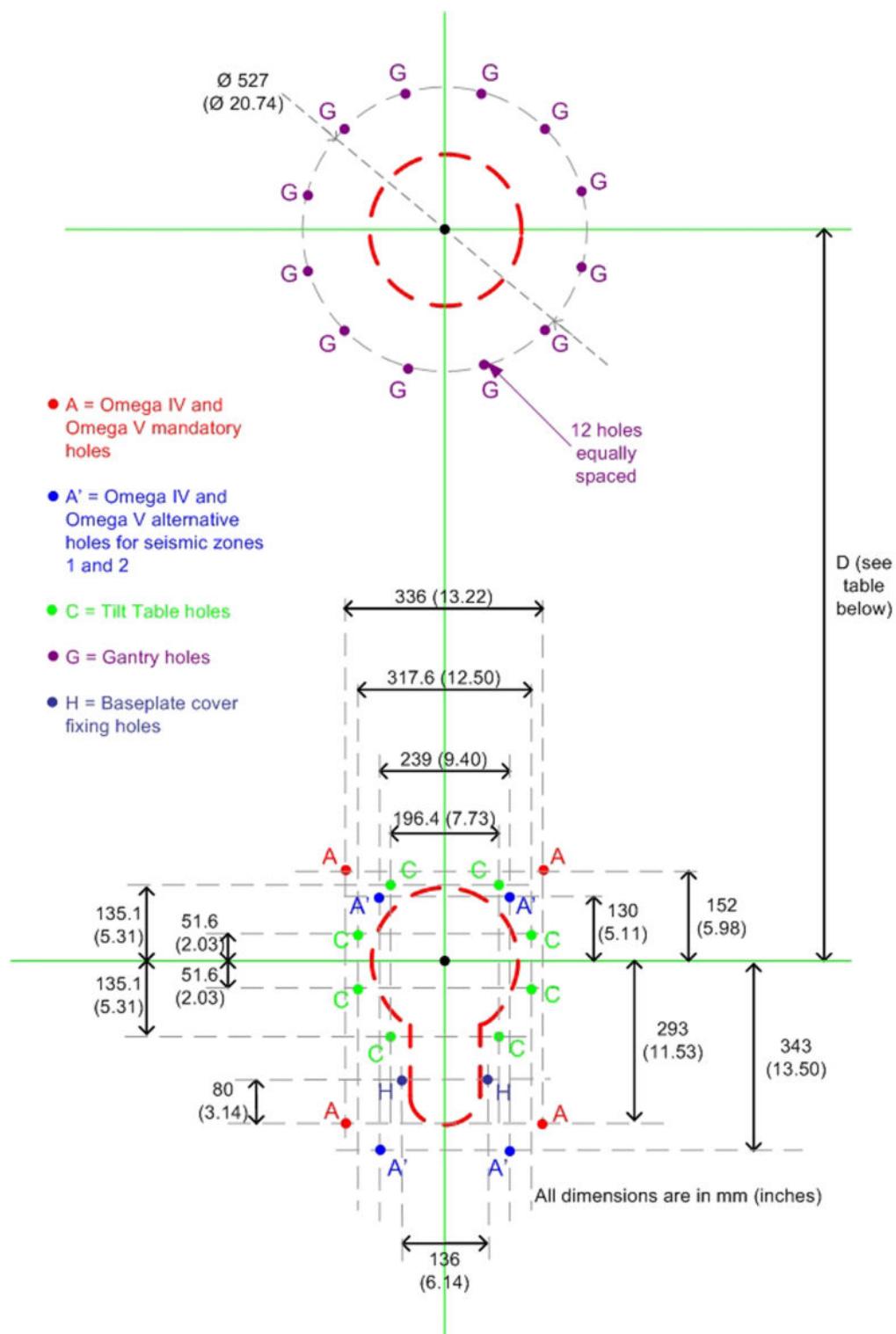
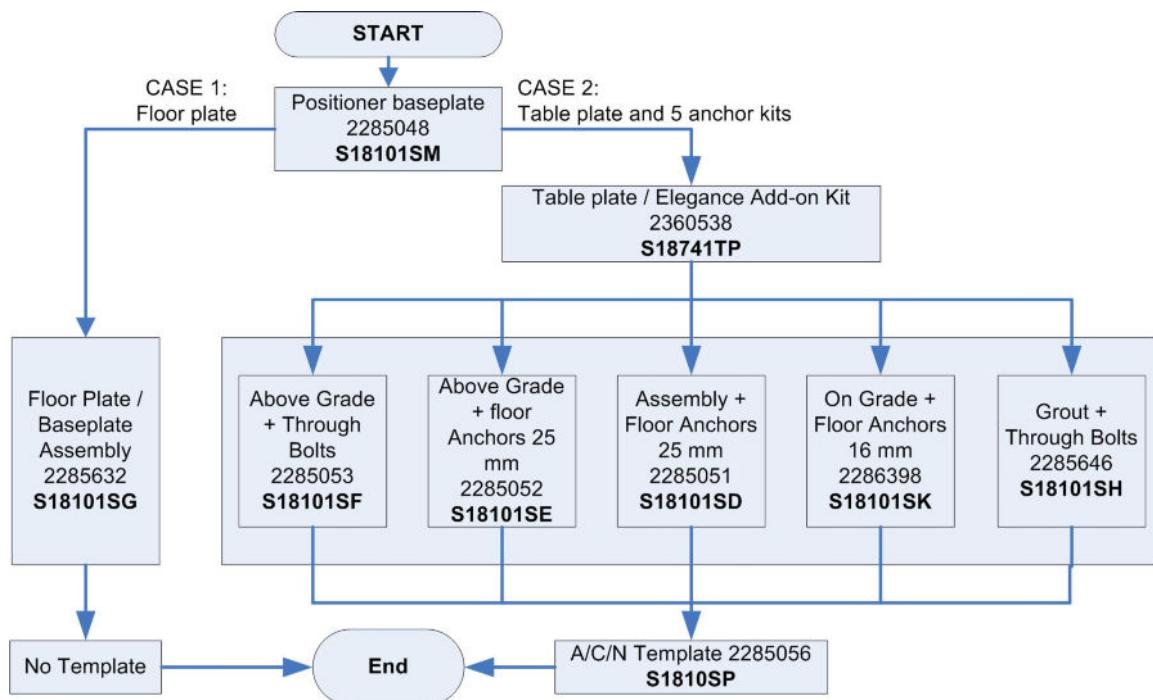


Table 2-11: D distance

	ANGIO	CARDIO	NEURO
Omega IV Compact	NA	1395 mm (54.9 in)	NA
Omega V Long	1278 mm (50.3 in)	1395 mm (54.9 in)	1395 mm (54.9 in)
Omega V non motorized Long	1278 mm (50.3 in)	1395 mm (54.9 in)	1395 mm (54.9 in)
Tilting table	1278 mm (50.3 in)	1395 mm (54.9 in)	1595 mm (62.8 in)

3.3.3 Gantry and Omega table/Tilting table Floor Preparation Kits (GEMS supplied)

Illustration 2-81:



All GE supplied vascular system floor preparation are contained in catalog. There are some additional gantry/table mounting kits based on each mounting method: through bolts or floor anchors.



NOTICE

The table base plate is recommended if a future Omega to Tilting table upgrade is to be considered.

- Base plate assembly (mandatory) 2285048 - Refer to [Table 2-12](#)
- Tilting table Add-On Kit 2360538 - (refer to [Table 2-13](#)) (Floor base plate assembly 2285632 shall be ordered).
- If there is no integrated base plate plan (2359194) delivered with S18101SG, please refer to drawings 2359194 (Angio and Cardio configuration [Illustration 2-82](#)) Please order locally the integrated base plate based on this plan.

- Assembly and separation (Select kit) 2285050 - Refer to tables, [Section 3.3.3.3](#)
 - Above grade, Floor anchors without insert – 2285051.
 - Through bolts without insert – 2285646.
 - On grade, Floor anchors – 2286398.
 - Above grade, Floor anchors and insert – 2285052.
 - Through bolts and insert – 2285053.
 - Floor plate / Base plate Assembly – 2285632.
- Templates (select kit) – 2285054 – Refer to [Section 3.3.3.4](#)
Angio/Cardio/Neuro templates – 2285056 (containing 2185979, 2127792 and 2360133 templates).

3.3.3.1 2285048 - Base Plate Assembly

Table 2-12:

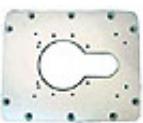
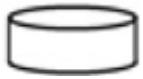
Item	Name	Part #	Description	Quan.	Notes
	Base plate	2285059	12 Hole Floor Base plate	1 pc	
	Lift Plate	2290939	Lifting Straps	2 pc	
	Hex Head Screw	5166535	Hex Head Screw 1/2"-13UNC L1 1/2" 12.9 Black	2 pc	
	Grease	2295599	Lithium grease lubricant 100g	2 pc	
	Doc	2230112-1-100	Vascular Gantry Base plate and Table	1 pc	
	Doc	2229297-100	LCA/LCV+/LC+ System Base plate and Omega IV/V/EP Table Floor Preparation	1 pc	
	Doc	2290880-2-100	Innova Pre-Installation Kit Install Procedure	1 pc	

NOTE: The 12 LC base plate mounting screws are delivered with the system's LC Gantry.
 These screws are:

- cap screws 5166774; Screw Socket Head Cap 1/2"-20 UNF L1 1/2" 12.9 Black (used when installing new system on old base plate (US threads))
- cap screws 2300939; Hex Head Cap screw; M12 40/40 Class 12.9 Black ; used to attach L-brackets on table shipping pallet to positioner dolly for table positioning (used when the new base plate is metric threads)

3.3.3.2 2360538 – Tilting table Add-On Kit

Table 2-13:

Item	Name	Part #	Description	Quan.	Notes
	Omega table/ Tilting table common Plate	2361993	Plate to be anchored under the table	1 pc	
	Hex Screws	5120708	Screw M16x40x40 Inox A4-70 Pass	10 pc	4 only are used for Omega
	Washer, Flat	99125091	Washer Plain - Large 17 mm/40 mm	10 pc	
	Floor Anchor	46-302265P1	5/8 diameter 6" floor anchor bolts	6 pc	
	Dowel	2290937-2	Wood Dowel; 16 mm diameter	6 pc	
	Bolt, Hex	2296892	Through bolt M20-500-400	6 pc	
	Washer, Flat	99142204	Washer plain 21 mm/40 mm for Through Bolts; one for each bolt	6 pc	
	Plate	2290941	Special Steel Spacer Plate; 4 in. x 4 in. (102 mm x 102 mm); one for each bolt	6 pc	
	Nut, Hex	99141607	Hex Nut M20 STL galvanized, two for each bolt	12 pc	
	Dowel	2290937	Wood Dowel; 24 mm diameter	6 pc	
	Cap	5130979	Plastic Cap	10 pc	
	M16 Plug	5130982	Plastic Plug	6 pc	see note *

NOTE: * M16 plug is needed only when Omega V table is replaced by Tilting table. Therefore, these plugs have to be kept (bag let inside the table base) until Tilting table is installed (no need to fit them with Omega table).

3.3.3.3 2285050 – Assembly and separation select kit

3.3.3.3.1 2285051 – Above grade, Floor Anchors without insert

Table 2-14:

Item	Name	Part #	Description	Quan.	Notes
	Floor Anchor	46-302265P1	Floor Anchor Bolt; 5/8 - 10x6" - (16 mm) drilling, (16 mm) drilling. 12 anchors for LC Positioner and 4 anchors for Table	16 pc	Above Grade anchor mounting method hardware
	Grout assy	2285055	- 10 kg Powdered Mortar Ardex K-15 - RTV Silicon Rubber Adhesive - 18 mm masking tape adhesive - Open cell foam	1 kit	Used in constructing LC Positioner grout dam
	Dowel	2290937-2	Wood Dowel; (16 mm) diameter	12 pc	

3.3.3.3.2 2285646 – Through bolts without insert

Table 2-15:

Item	Name	Part #	Description	Quan.	Notes
	Bolt, Hex	2296892	Through Bolt; M20 - 500-400 12 bolts for LC Positioner and 4 bolts for Table	16 pc	Through Bolt mounting method hardware
	Washer, Flat	99142204	Special Flat Washer for Through Bolts; one for each bolt	16 pc	same as above
	Plate	2290941	Special Steel Spacer Plate; 4 in. x 4 in. (102 mm x 102 mm); one for each bolt	16 pc	same as above
	Nut, Hex	99141607	Hex Nut M20 STL galvanized two for each bolt	32 pc	same as above
	Grout assy	2285055	- 10 kg Powdered Mortar Ardex K-15 - RTV Silicon Rubber Adhesive - 18 mm masking tape adhesive - Open cell foam	1 kit	Used in constructing LC Positioner grout dam
	Dowel	2290937	Wood Dowel; 24 mm diameter	12 pc	

3.3.3.3.3 2286398 – On grade, Floor anchor

Table 2-16:

Item	Name	Part #	Description	Quan.	Notes
	Floor An-chor	46-302265P1	Floor Anchor Bolt; 5/8-10 x 6 in. 12 anchors for LC Positioner and 4 anchors for Table	16 pc	On Grade anchor mounting method hardware
	Grout assy	2285055	- 10 kg Powdered Mortar Ardex K-15 - RTV Silicon Rubber Adhesive - 18 mm masking tape adhesive - Open cell foam	1 kit	Used in constructing LC Positioner grout dam
	Dowel	2290937-2	Wood Dowel; 16 mm diameter	12 pc	
	Cable Con-duit	2285057	Cable conduit - sheet metal part	1 pc	
	Vinyl Trim	2296890	Gripping Range; 1.5 to 3 mm THK	1 m	
	Vinyl Trim	2296891	Vinyl trim with segmented metal core 12 mm	1 m	

3.3.3.3.4 2285052 – Above grade, Floor anchor with insert

Table 2-17:

Item	Name	Part #	Description	Quan.	Notes
	Floor An-chor	46-302265P1	Floor Anchor Bolt; 5/8-10 x 6 in. drilling 12 anchors for LC Positioner and 4 anchors for Table	16 pc	Above Grade anchor mounting method hardware
	Grout assy	2285055	- 10 kg Powdered Mortar Ardex K-15 - RTV Silicon Rubber Adhesive - 18 mm masking tape adhesive - Open cell foam	1 kit	Used in constructing LC Positioner grout dam
	Dowel	2290937-2	Wood Dowel; 16 mm diameter	12 pc	
	Water Elec-tric Sepa-rator	2268647	Manufactured part introduced in Positioner key hole to fit 1 electrical conduit of 6".	1 pc	Electrical and water conduits separated inch size.

3.3.3.3.5 2285053 – Through bolts with insert

Table 2-18:

Item	Name	Part #	Description	Quan.	Notes
	Bolt, Hex	2296892	Through Bolt; M20 - 500-400 12 bolts for LC Positioner and 4 bolts for Table	16 pc	Through Bolt mounting method hardware
	Washer, Flat	99142204	Special Flat Washer for Through Bolts; one for each bolt	16 pc	same as above
	Plate	2290941	Special Steel Spacer Plate; 4 in. x 4 in. (102 mm x 102 mm); one for each bolt	16 pc	same as above
	Nut, Hex	99141607	Hex Nut M20 STL galvanized two for each bolt	32 pc	same as above
	Grout assy	2285055	- 10 kg Powdered Mortar Ardex K-15 - RTV Silicon Rubber Adhesive - 18 mm masking tape adhesive - Open cell foam	1 kit	Used in constructing LC Positioner grout dam
	Dowel	2290937	Wood Dowel; 24 mm diameter	12 pc	
	Water Elec- tric Sepa- tor	2268647	Manufactured part introduced in Positioner key hole to fit 1 electrical conduit of 6".	1 pc	Electrical and water con- duits separated inch size.

3.3.3.3.6 2285632 – Floor plate / Base plate assembly

Table 2-19:

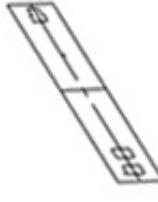
Item	Name	Part #	Description	Quan.	Notes
	Hex Screws	2360523	Screw M16x30x30 Inox A4-80 Pass	12 pc	For floor plate / Base plate assembly
	Washer	99125091	Washer P 17 mm/40 mm	34 pc	24 are used for the positioner integration
	Hex Screws	99133570	Screw M16x40x40 Inox A4-80 Pass	10 pc	4 only are used for Omega

NOTE: The kit 2285632 Floor plate / Base plate assembly is needed when the floor plate is installed. This part isn't GE part. It's ordered locally.

3.3.3.4 2285054 – Templates

3.3.3.4.1 2285056 - A/C/N Templates

Table 2-20:

Item	Name	Part #	Description	Quan.	Notes
	A/C/N Tem-plates	2185979	LC Positioner and Omega C/N Pa-tient Table floor mounting tem-plate	1 pc	See Illustration 2-80
		2127792	LCV+ Gantry and Omega A Pa-tient Table floor mounting tem-plate	1 pc	
		2360133	Innova 4100 Template	1 pc	Specific for System with 41 cm detector Tilting table template

3.3.3.4.2 2186109 – EP Template

Table 2-21:

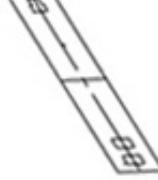
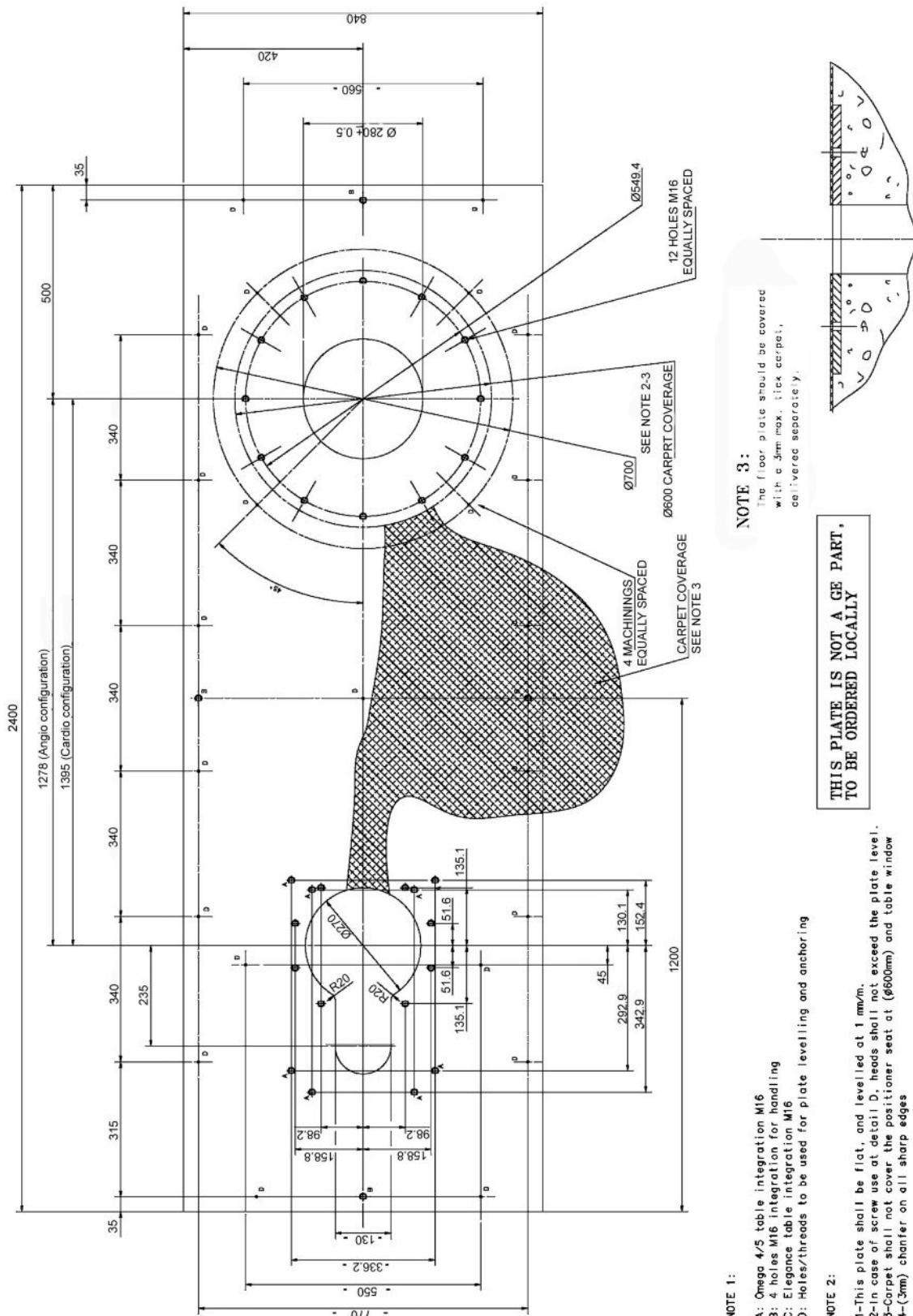
Item	Name	Part #	Description	Quan.	Notes
	EP Tem-plate	2186109	LCV+ Gantry and EP Patient Ta-ble floor mounting template	1 pc	

Illustration 2-82: Base plate plan 2359194 - Angio and Cardio configurations



3.3.4 Injector Mounting Requirements



CAUTION

Table accessory rail load consideration:

The maximum load per table accessory rail is 40 kg (88 lbs) at 150 mm (0.49 ft) (60 N.m or 44.25 ft/lbs). Therefore:

- Only light extra load not exceeding 5 kg (11 lbs) at 100 mm (0.33 ft) (i.e IV pole with its accessories, pressure head...) is authorized on the same table accessory rail as the injector.
- Never install injector and radiation protection on the same table accessory rail.
- Typical installation on the front table accessory rail is Smart handle or Smart box, Table Side System Control (TSSC), Innova Central/Touchscreen, Table panning device and cables support.
- If needed an optional rail can be installed at table foot end of the Omega V table for other options.

3.3.5 Prerequisites for ECG Acquisition Kit Installation

3.3.5.1 General Prerequisites for ECG Acquisition Kit Installation

Verify the following items to ensure easy mounting for the Hubican & Physio modules in all situations:

1. Check enough space can be managed to properly install the Hubican module in Control Room.
2. Check enough space can be managed to route/hide Hubican cable (VCIM-to-Hubican 12m long cable) in Control Room.

3.3.5.2 Specific Prerequisites for Installation Configuration #1

Verify the following items only in the case of Installation Configuration #1 with ECG Device in Control Room:

1. Check the environment of the ECG Device module located in Control Room provides enough space to install the Physio module at a max distance of 1 meter.
2. Check enough space can be managed to properly route/hide Physio cable (Physio-to-Hubican) in the Control Room
3. The Analog Output Box option is mandatory to provide on analog output connection to the Physio module (If not present, it can be ordered through the following FRUs):
 - 2018971-001 16CH ANALOG OUTPUT CPU INTERFACE OPTION
 - 2007557-002 KIT ANALOG OUTPUT BOX W/CABLES
 - 2010476-001 BOX CARDIOLAB/MACLAB ANALOG OUTPUT

3.3.5.3 Specific Prerequisites for Installation Configuration #2

Verify the following items only in the case of Installation Configuration #2 with ECG Device in Exam Room:

1. Check the environment of the ECG Device module located in Exam Room provides enough space to install the physio module at a max distance of 1 meter.
2. Check enough space can be managed to properly route/hide Physio cable (Physio-to-Hubican) at table base area.
3. Check the conduit normally designed for MacLab signal cables is present (see conduit illustration in [Chapter 5, Cable Channeling](#)).

3.3.6 Large Display Subsystem option



NOTICE

General safety instructions

- Move the LD Cabinet & LD Monitor in an upright position in their original packages to the final destination room. To lift the LD cabinet, use a forklift or lifting belts with spreader bars.
- Check for sufficient floor and elevator loading capacity.
- Check the integrity of the LD Subsystem equipment carefully.
- If you notice visible damage, do not install or start the LD Subsystem equipment. Contact the nearest Service Center immediately.
- All installation, maintenance and service work should be performed by qualified Service personnel.

3.4 Ceiling Requirements

Aluminum rails support the In-Room Monitor bridge used in Innova system X-Ray rooms.

Reference:

For additional details on ceiling requirements for stationary rails, refer to: - Direction 46-019639, *Advantx (VHLA) XT Stationary Rails Installation and Adjustment*.

When evaluating ceiling you must take into account the following mounting information:

3.4.1 Rail Mounting

Attach stationary rails to structural steel with through-bolts in concrete ceilings. Do not use screw anchors in direct tension.

Mount stationary rails directly to the ceiling slab or to flush-mounted unistrut or halfen structure. In higher rooms with false ceiling, mount stationary rails to rigid vertical members hung from ceiling slab.

Securing a supplementary channel to the bottom of the vertical members and mounting the stationary rails to this channel can greatly reduce the number of vertical members.

The stationary rail support structure must be leveled before installation can begin. Do not assume that any support structure is level within specified tolerances, particularly after removing suspensions from an existing room.

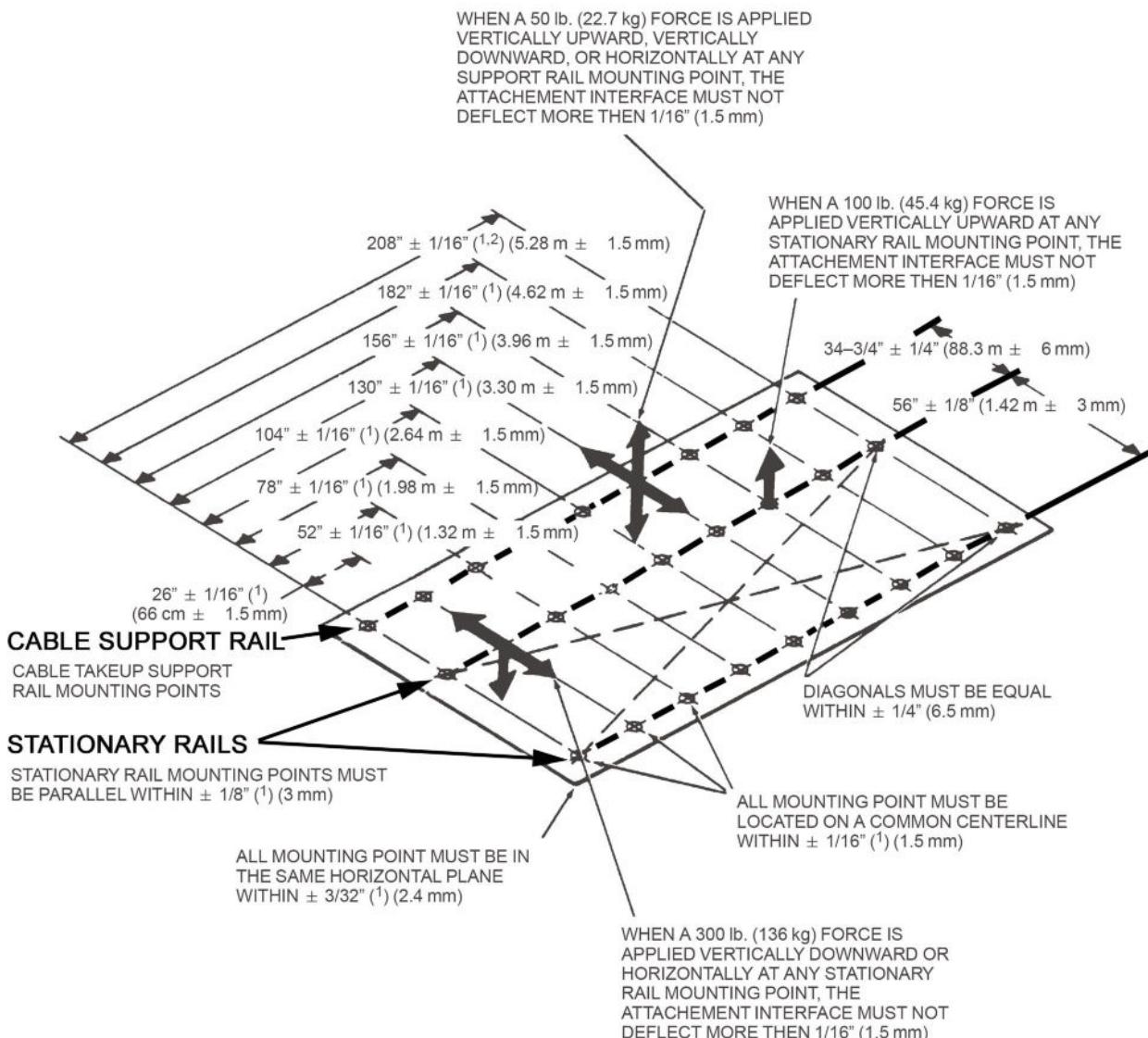
3.4.2 Bolt Specifications (Mavig suspensions)

- The maximum load per bolt will not exceed **1557 N**.
- Each bolt must not “pull out” or otherwise fail under a vertically downward *dead* load of **6227 N**.

3.4.3 Select Rails (Mavig suspensions)

All XT Stationary rails are with a select length process. Detail of available length is illustrated in [Chapter 1, European Process Order Select](#).

Illustration 2-83: SPECIFICATIONS FOR A TYPICAL 17'-10" (5.44 M) INBOARD STATIONARY RAIL MOUNTING INTERFACE (BOTH RAILS CEILING MOUNTED), FOR MAVIG SUSPENSION



NOTES: 1. NONE CUMULATIVE ERROR.
2. SPACE BETWEEN LAST 2 HOLES MAY BE LESS THAN 26" (66 cm)

Table 2-22: Stationary rail in different length

Rail length mm (ft)	A	C	D	INBOARD RAILS
4,724 (15'6")	7*660.4=4,623		51	B0186JA
5,639 (18'6")	8*660.4=5,283	254	51	B0222JA
5,791 (19')	8*660.4=5,283	406	51	B0228JA

3.4.4 Boom Mounted for Monitor Suspension

The purchaser/customer should prepare a Unistrut/ Halfen or equivalent ceiling structure. The distance between center lines of the two longitudinal rails shall be 660.4 mm (26").

One type of boom mounted is supplied as counterpoised monitor suspension not adjusted on site as follows:

- for 2 x 21" (53 cm) EIZO GmbH (formerly Siemens) flat panel 21" (Hi-bright) and additional AW NEC monitor.

3.4.5 Cable Support for Monitor Cables

A cable support (cable drape) is provided with an Innova System.

The cable support kit contains:

- 1 B2054 EK (Drape with 3 M Bridge, on suspensions for X-Ray tubes and monitors, contains 8 FT 6 inch track, three carriers, and mounting hardware)
- 1 B2055 ED (stationary rail parts)

NOTE: In Americas the Cable Support Kit must be provided locally by the Customer (e.g. CPGE55 from Unistrut).

3.5 Wall Requirements

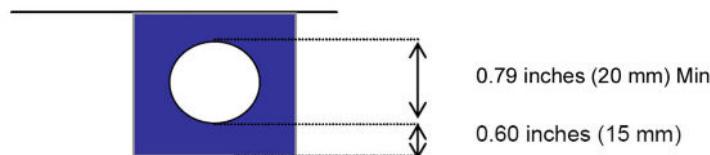
The C1 Cabinet, the C2 Cabinet, the PDB Cabinet and LD cabinet (optional) must be securely fastened to the wall to prevent them from tipping.

The optional LD secondary monitor outside the patient room is mounted on wall. The swingout arm that holds the LD additional monitor shall be mount according to the manufacturer mounting manual, see *Articulating Arm Wall Mount Installation Manual* in OEM manuals list.

An hooking point shall be provided in order to uplift the LD additional monitor at the swingout arm level during its installation:

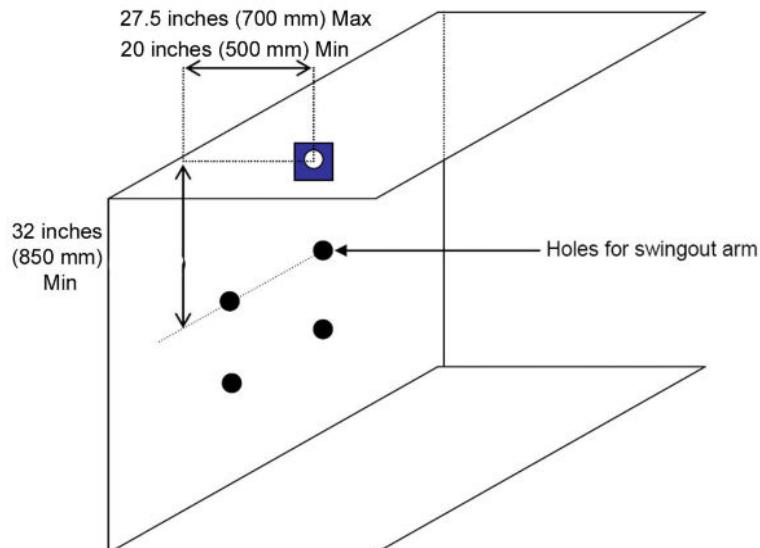
- Hooking point characteristic: It must withstand up to 440 lbs (200 Kgs)
- Recommended hooking point dimensions:

Illustration 2-84: hooking point dimensions



- Hooking point position:

Illustration 2-85: Hooking point position



4 Mounting Data, Including Seismic

4.1 Seismic areas

4.1.1 Floors

Chiller: The seismic kit for the Chiller 4100 (SMC) is included with the chiller when shipped.

LD cabinet: the seismic kit to fasten the LD cabinet to the wall is included with the cabinet when shipped.

In Seismic areas all cabinets must be anchored to the floor. See [Basic Innova System Compatibility](#) for referential documents. C1 & C2 Cabinets: Wall support 2284542-2.

Every sub-system is delivered on site with its proper seismic kit.

- Monitor Flat Panel Seismic Kit: 2353317
- VCIM seismic kit: 2365510.

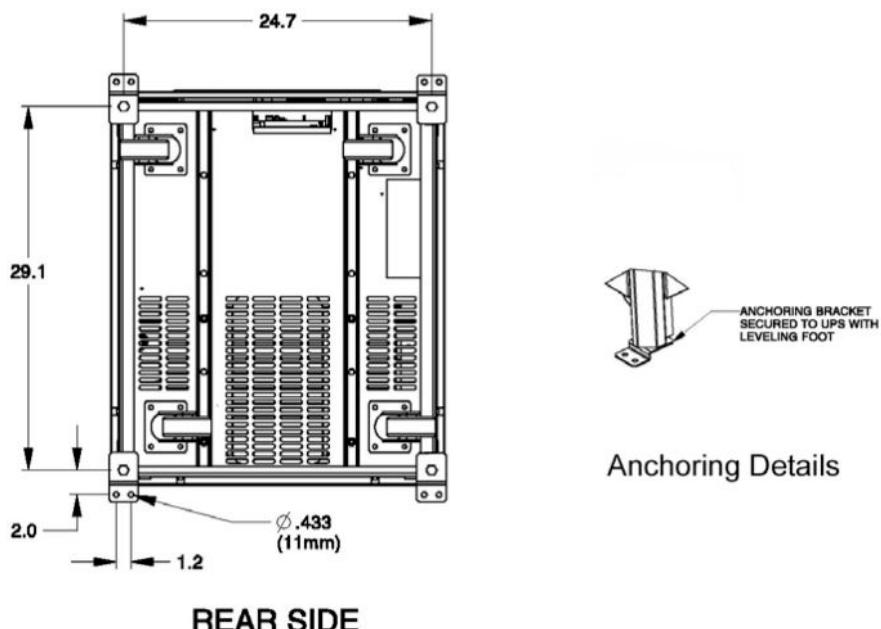
Anti-seismic means be installed before opening the system for normal use.

4.1.2 Walls

NOTE: For Fluoro UPS option (US version only), the Fluoro UPS supplier will deliver anchoring brackets (see [Illustration 2-86](#)). The bolts will be provided by the customer.

Consider local seismic codes when planning cabinet mounting. Consult seismic expert to determine which mounting method is appropriate for the seismic region. Certain seismic regions require additional reinforcement in walls. See [Basic Innova System Compatability](#) for referential documents.

Illustration 2-86:



4.2 Seismic Calculations

Seismic requirements are determined and specified by the hospital/ Design Professional of record and may require approval by the specific state or country agency.

Seismic attachment hardware shown on seismic calculations may differ from hardware supplied with system. Any additional hardware that is required will be the responsibility of the institution and/or their contractor. Contact your local GE Installation Program Manager to obtain seismic calculations.

Seismic calculations are per California Building Code (CBC) and International Building Code (IBC).

The following shows center-of-gravity information for system components:

- C1 cabinet, [Illustration 2-87](#)
- C2 cabinet, [Illustration 2-88](#)
- Fluoro UPS UL, [Illustration 2-89](#)
- Fluoro UPS CE, [Illustration 2-90](#)
- PDB UL, [Illustration 2-91](#)
- PDB CE, [Illustration 2-92](#)
- Tube Chiller Coolix 4100, [Illustration 2-93](#)
- Detector Conditioner, [Illustration 2-94](#)
- Omega IV Table, [Illustration 2-95](#)
- Omega V Long Table, [Illustration 2-96](#)
- Tilting table, [Illustration 2-97](#)
- Gantry, [Illustration 2-98](#)
- 3 kVa UPS (LD option), [Illustration 2-99](#)
- 19" Desk Mounted Monitor, [Illustration 2-100](#)
- Mavig Overhead Flat Panel Suspension, [Illustration 2-101](#)
- Large Display Cabinet (LD option), [Illustration 2-102](#)
- Large Display Monitor suspension (LD option), [Illustration 2-103](#)
- Large Display secondary monitor Swing out arm (LD option), [Illustration 2-104](#)

Illustration 2-87: C1 cabinet

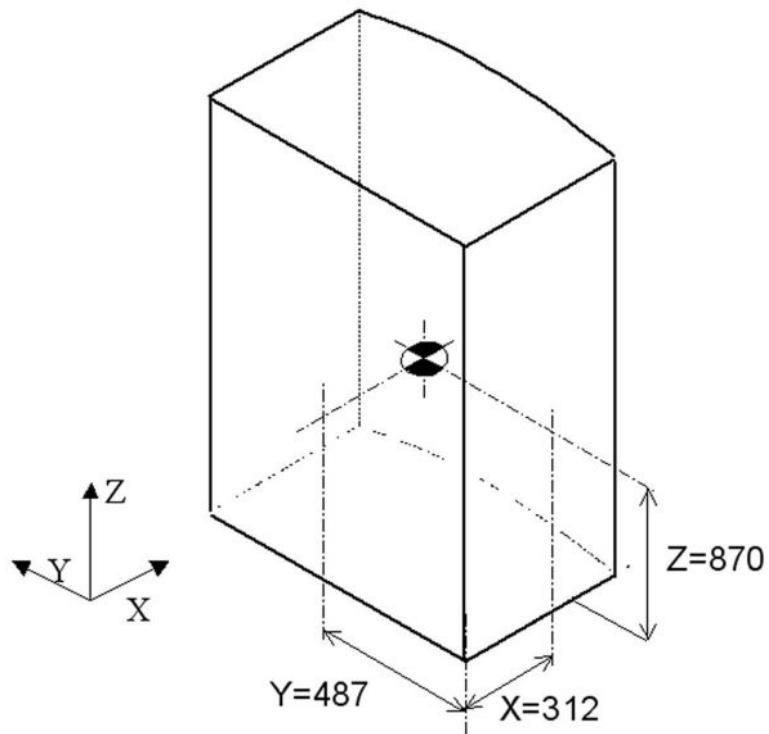


Illustration 2-88: C2 cabinet

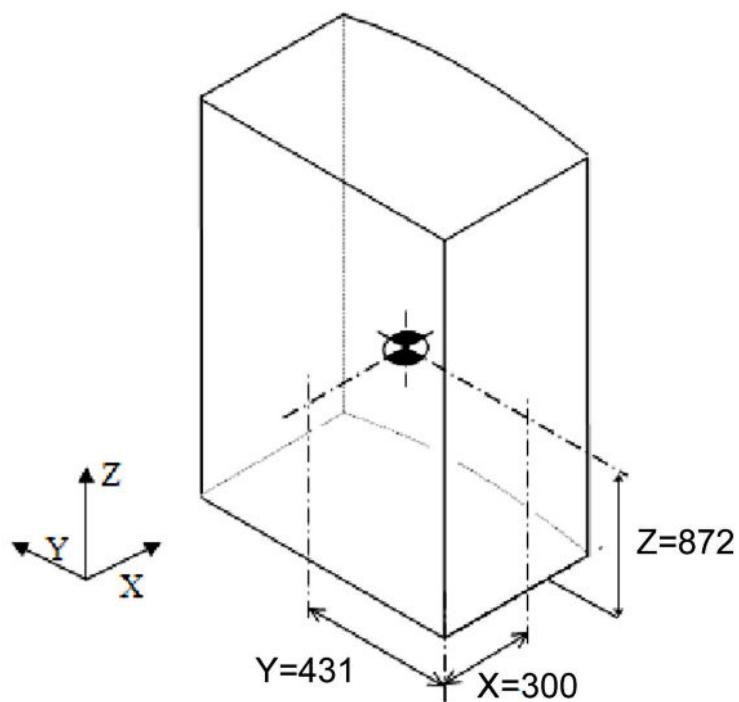


Illustration 2-89: Fluoro UPS UL

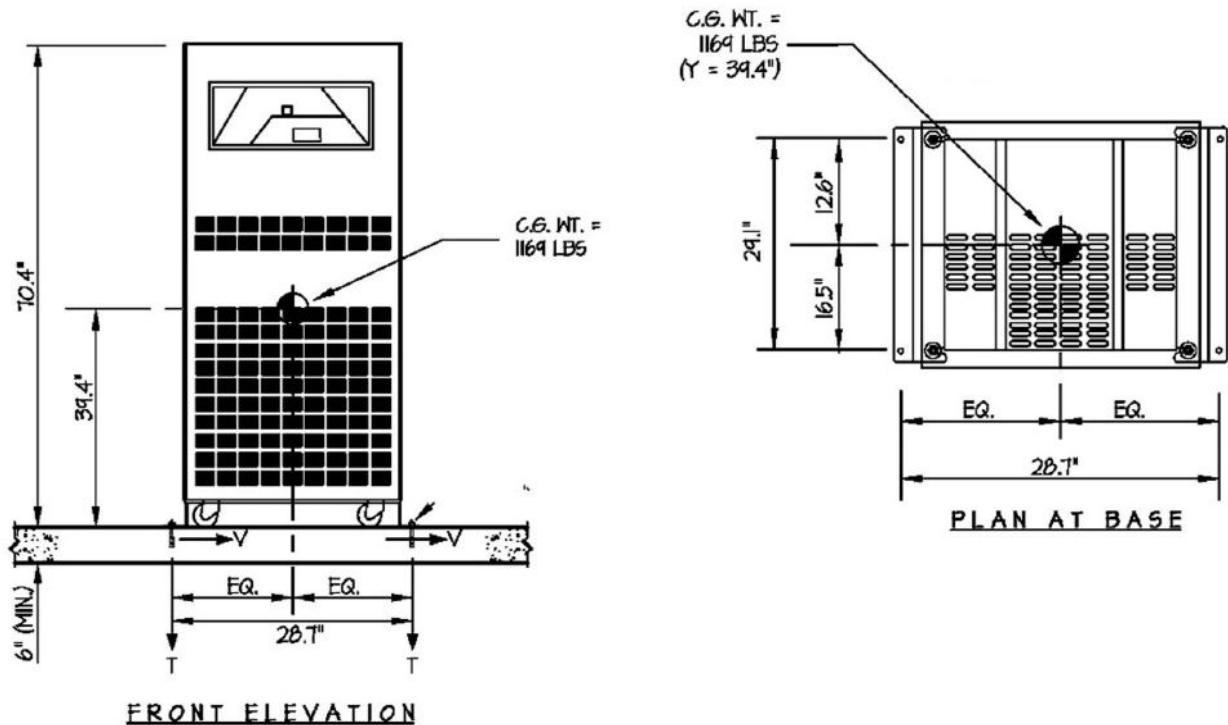


Illustration 2-90: Fluoro UPS CE

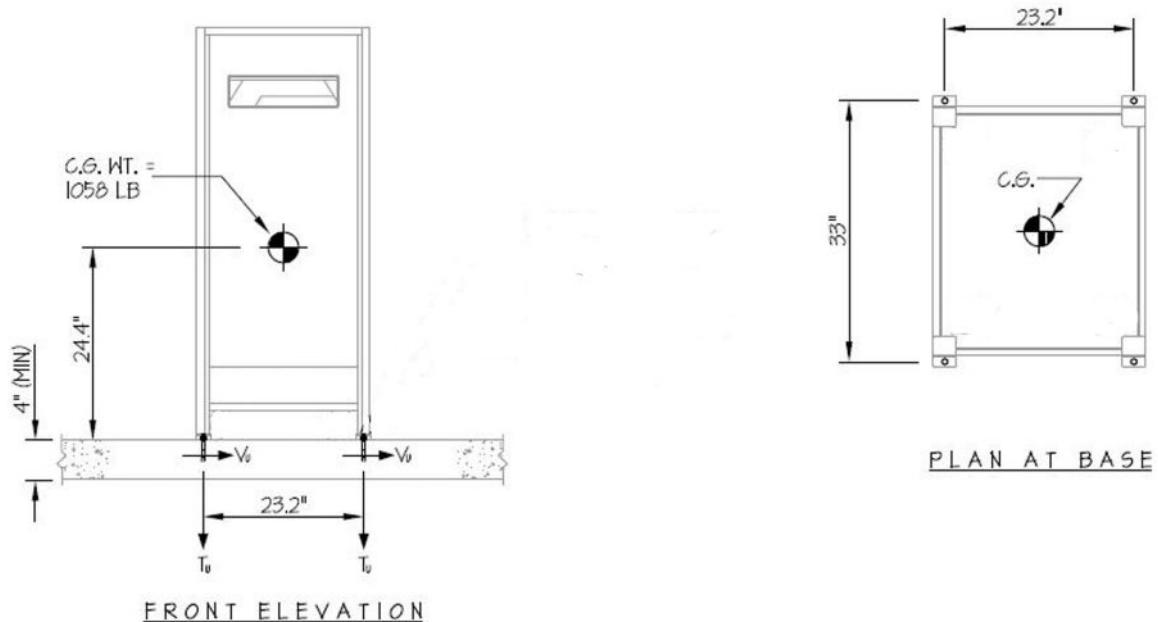


Illustration 2-91: PDB UL

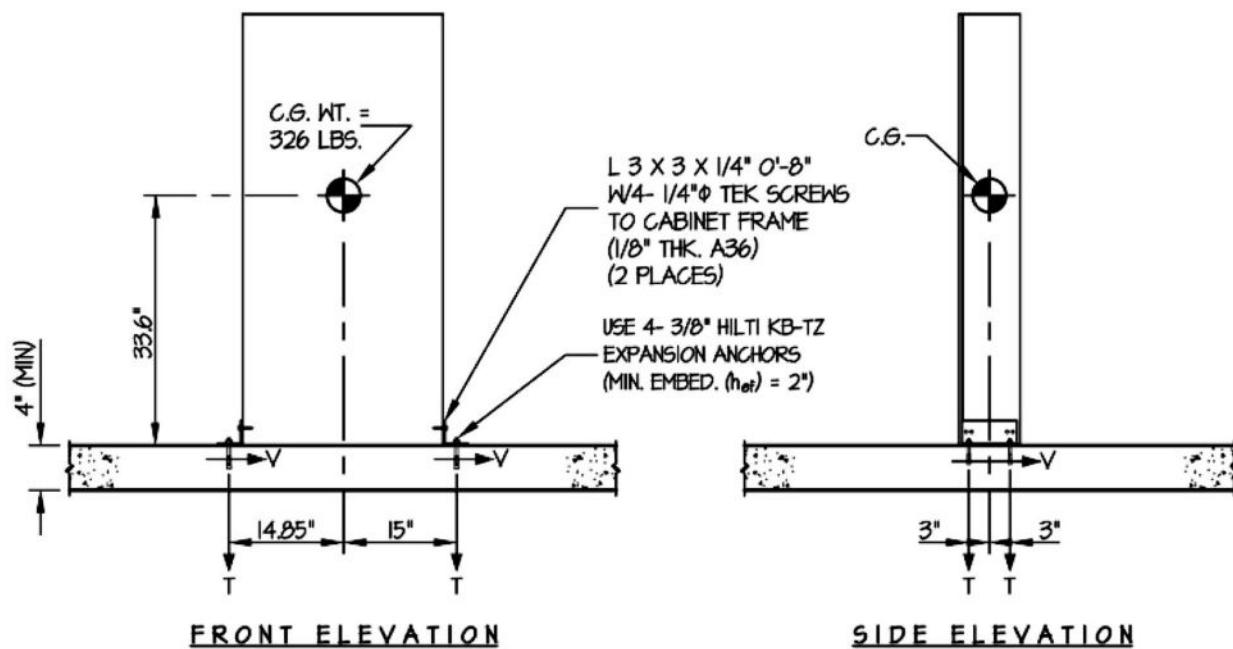


Illustration 2-92: PDB CE

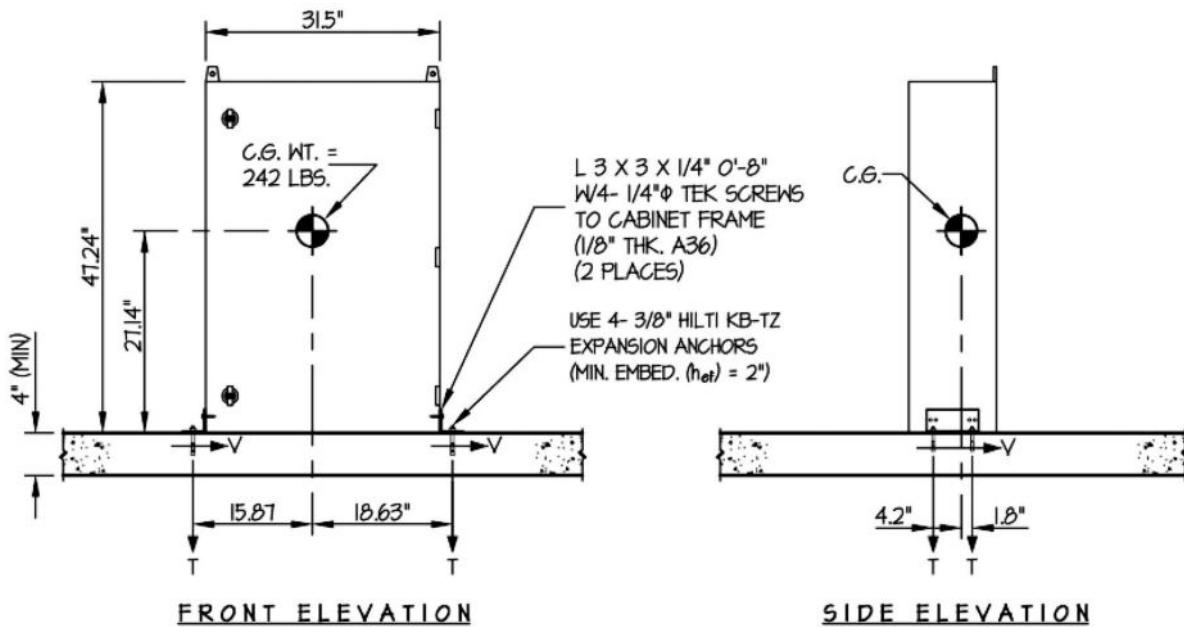


Illustration 2-93: Tube Chiller Coolix 4100

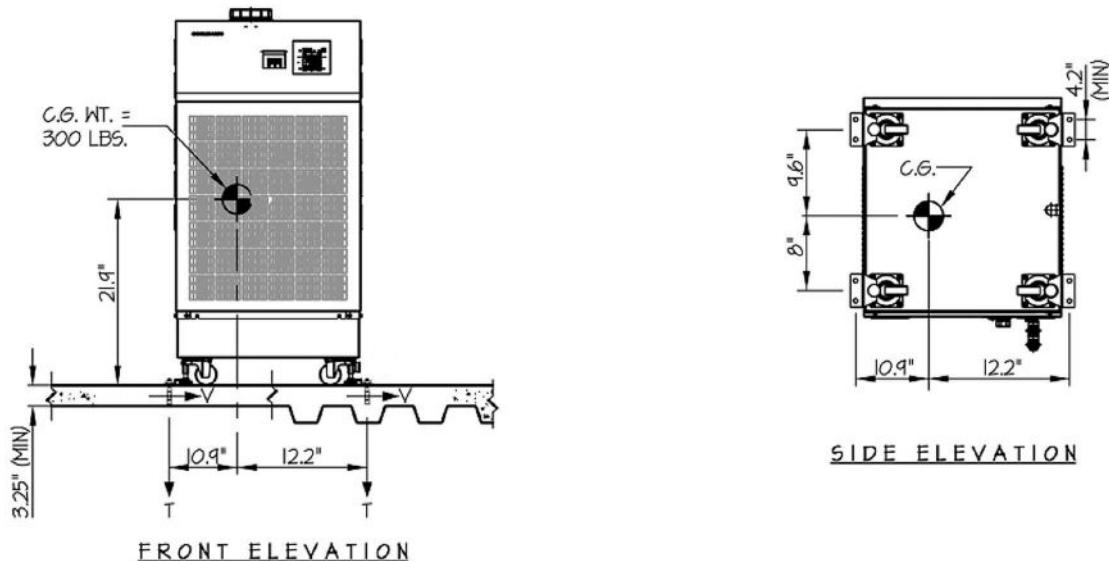


Illustration 2-94: Detector Conditioner

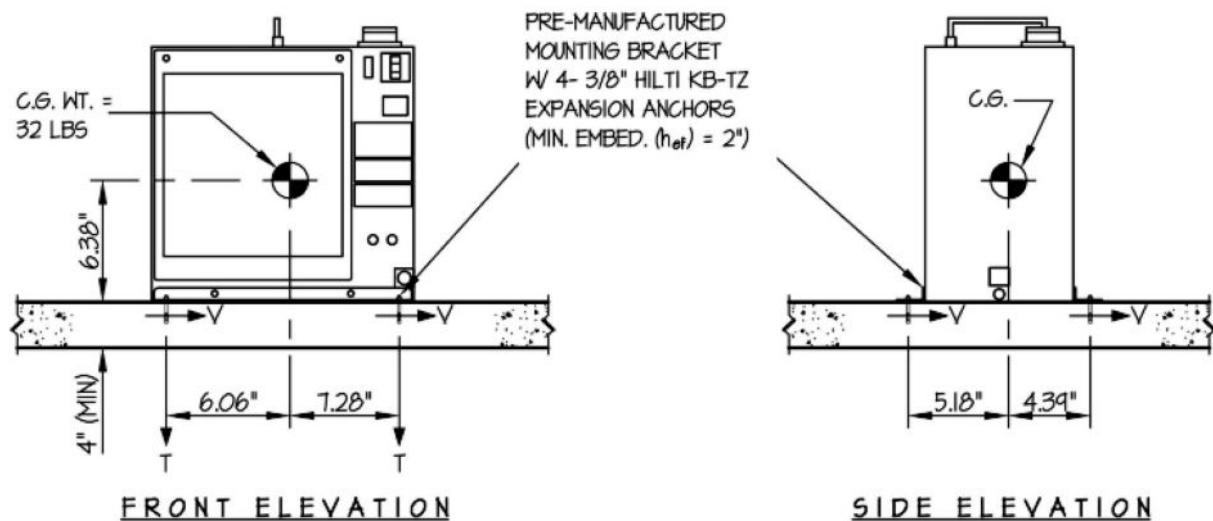


Illustration 2-95: Omega IV Table

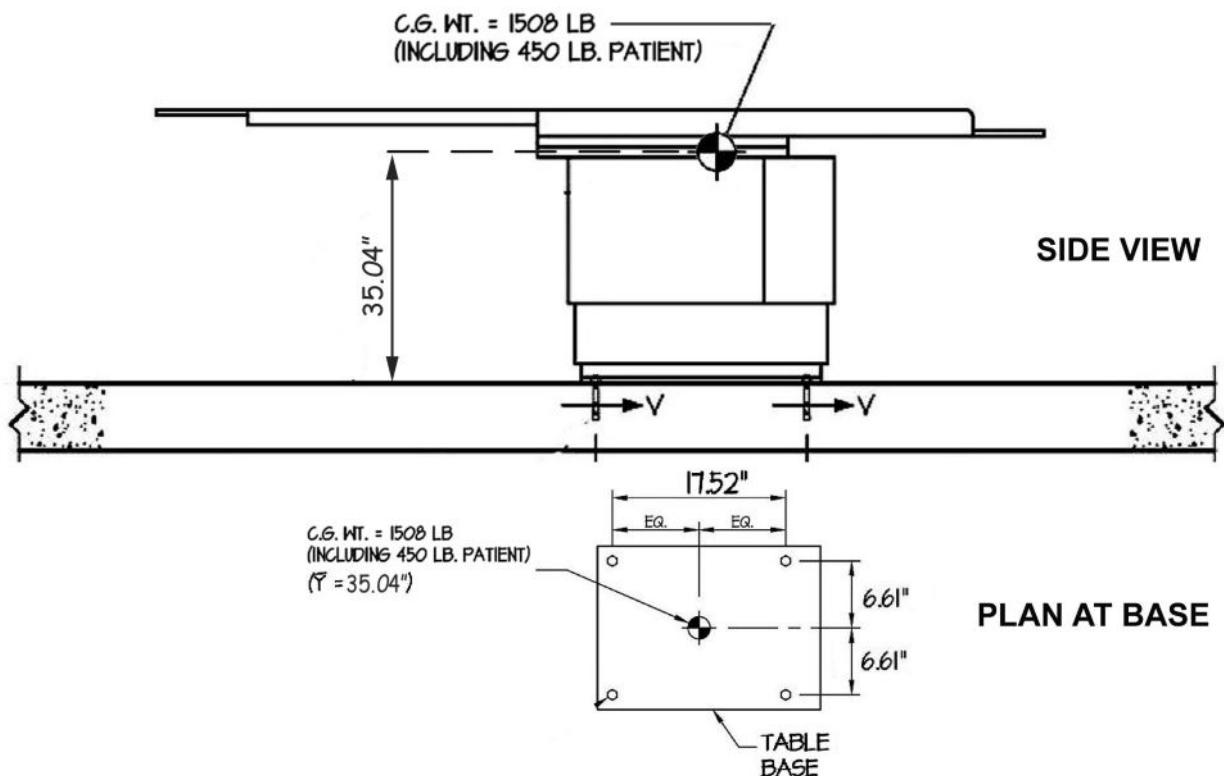


Illustration 2-96: Omega V Long Table

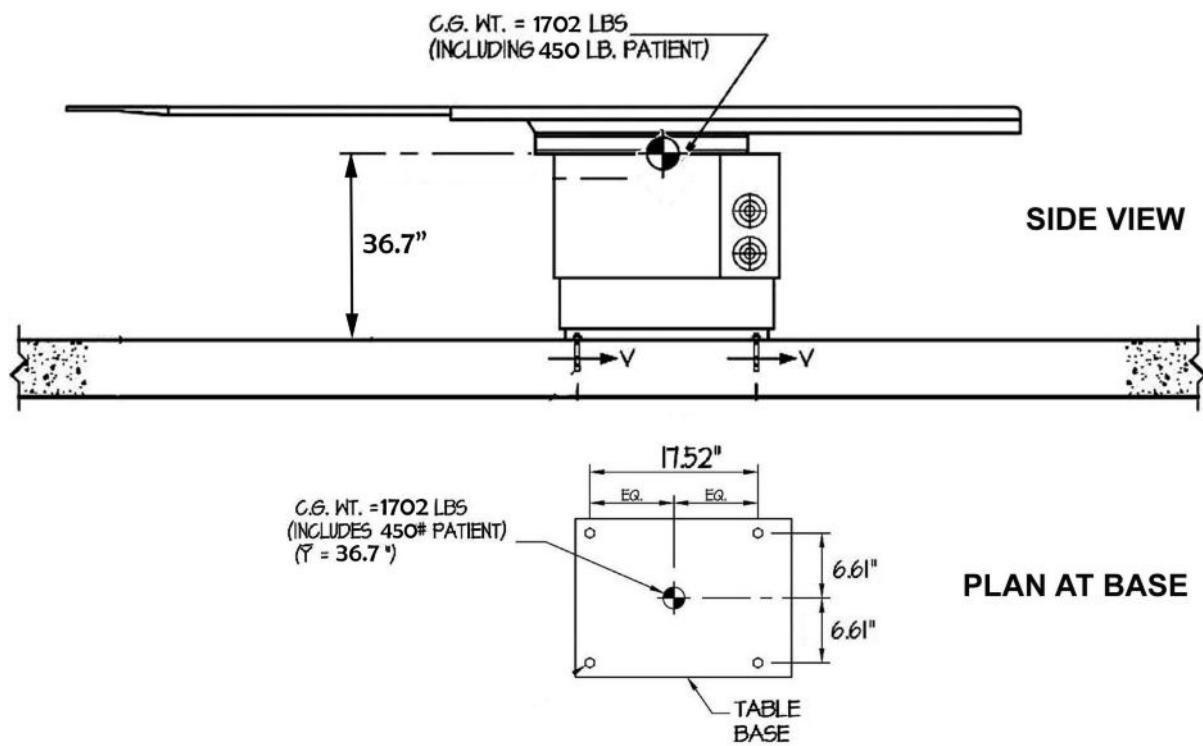


Illustration 2-97: Tilting table

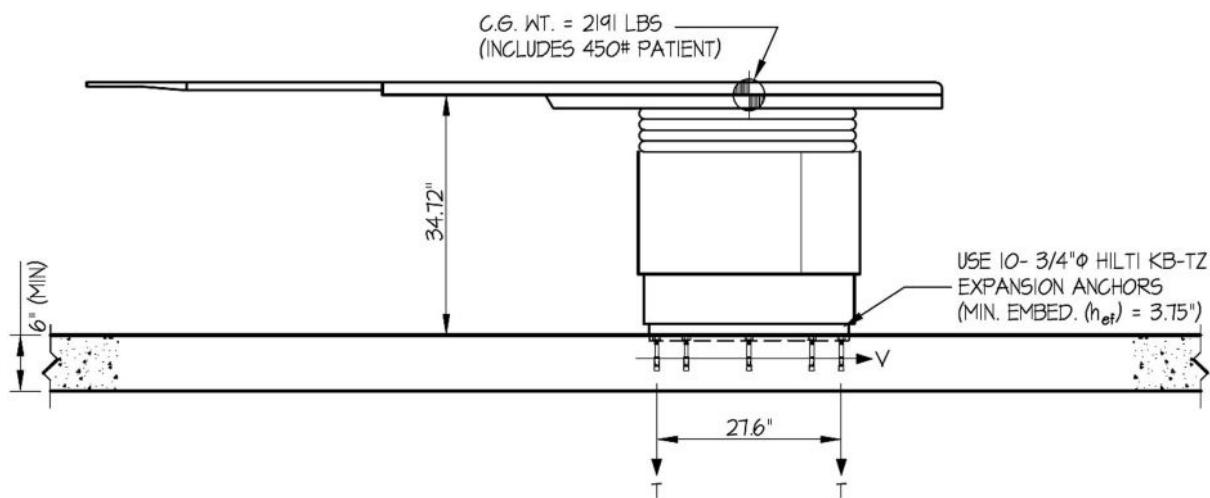


Illustration 2-98: Gantry

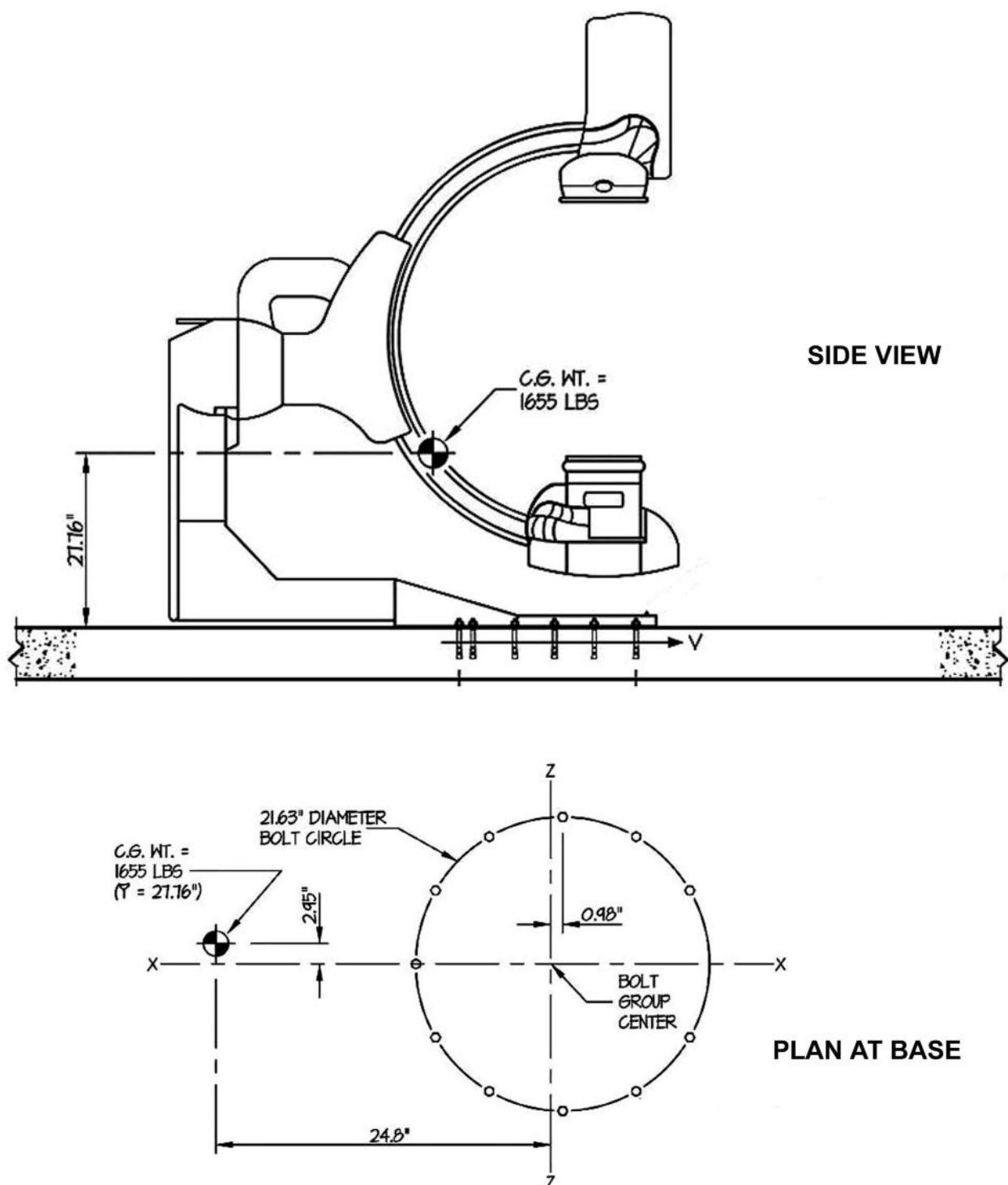


Illustration 2-99: 3 kVA UPS (LD option)

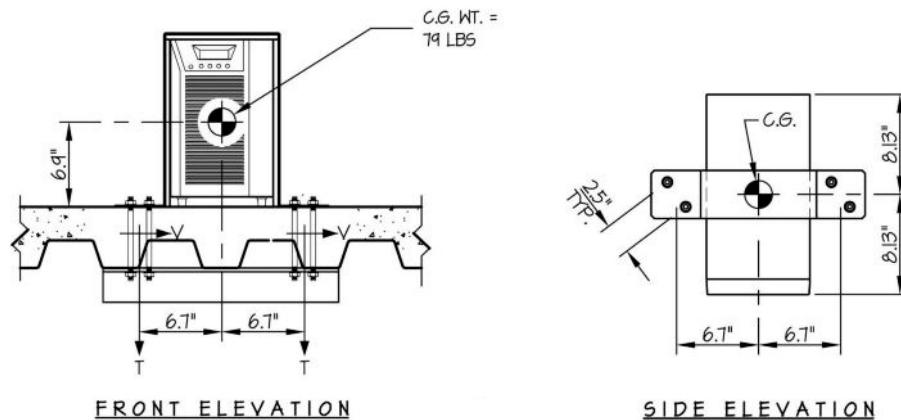


Illustration 2-100: 19" Desk Mounted Monitor

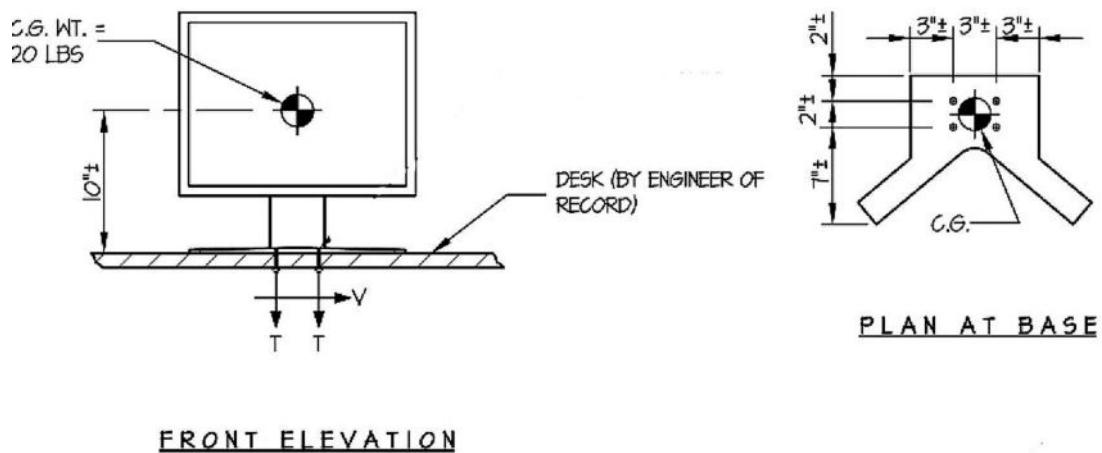


Illustration 2-101: Mavig Overhead Flat Panel Suspension

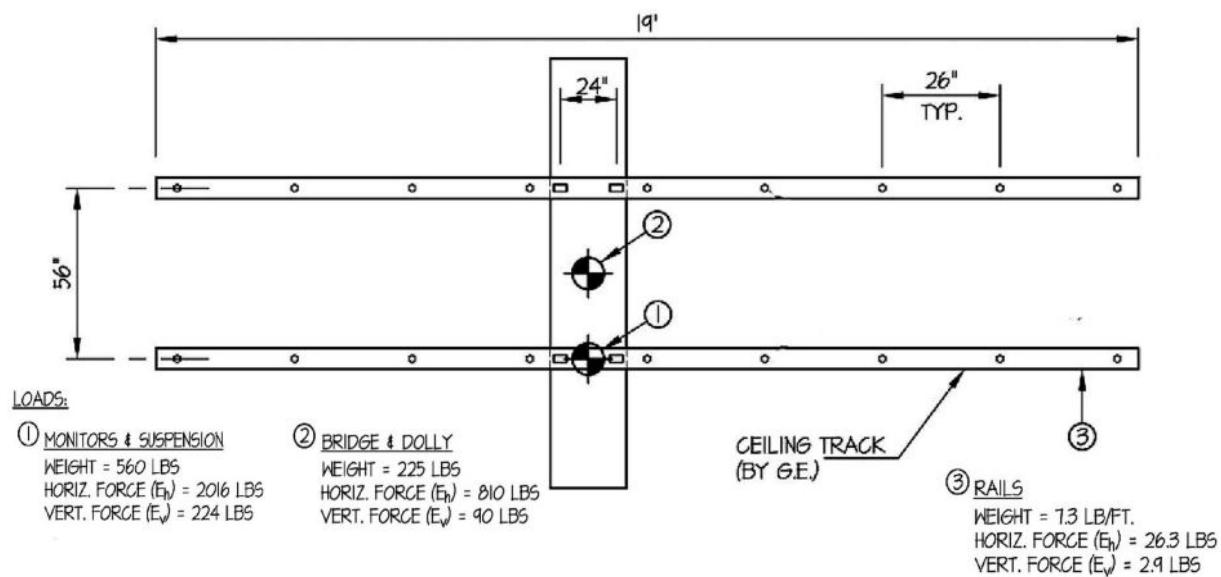
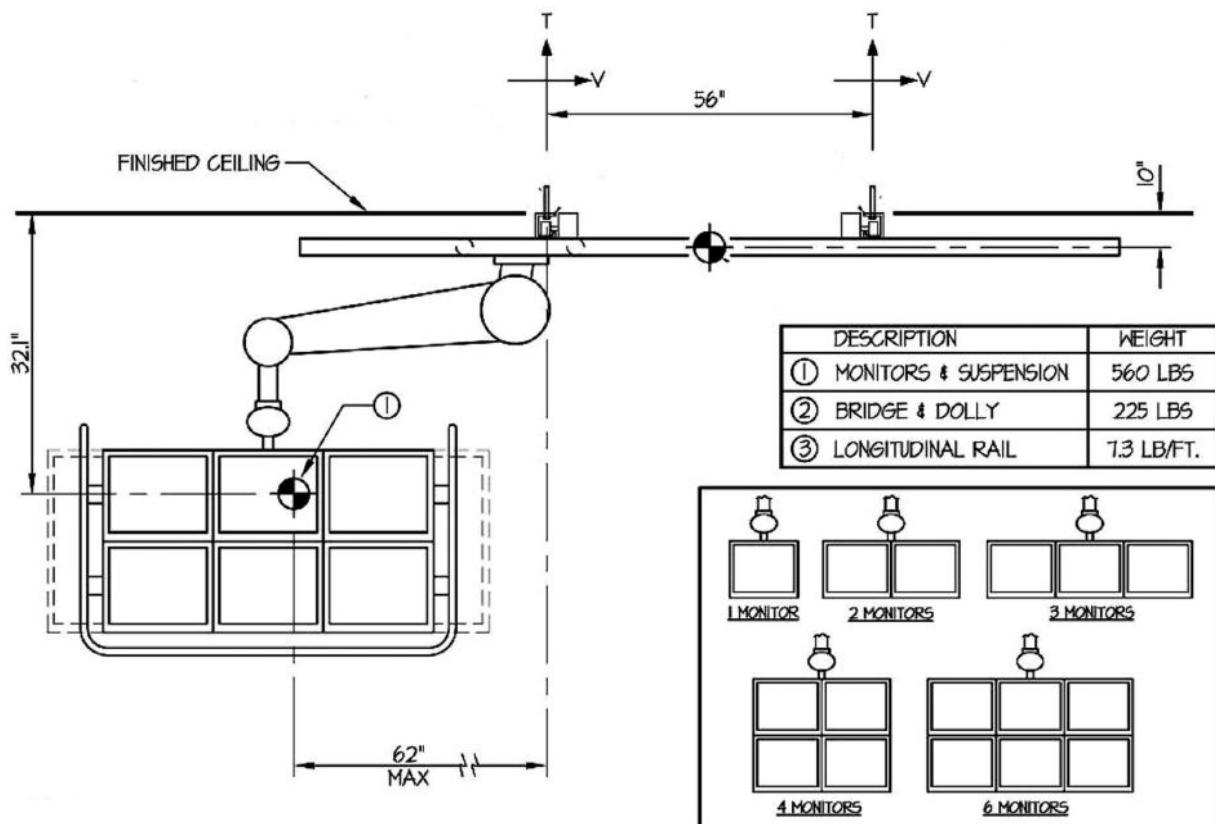


Illustration 2-102: Large Display Cabinet (LD option)

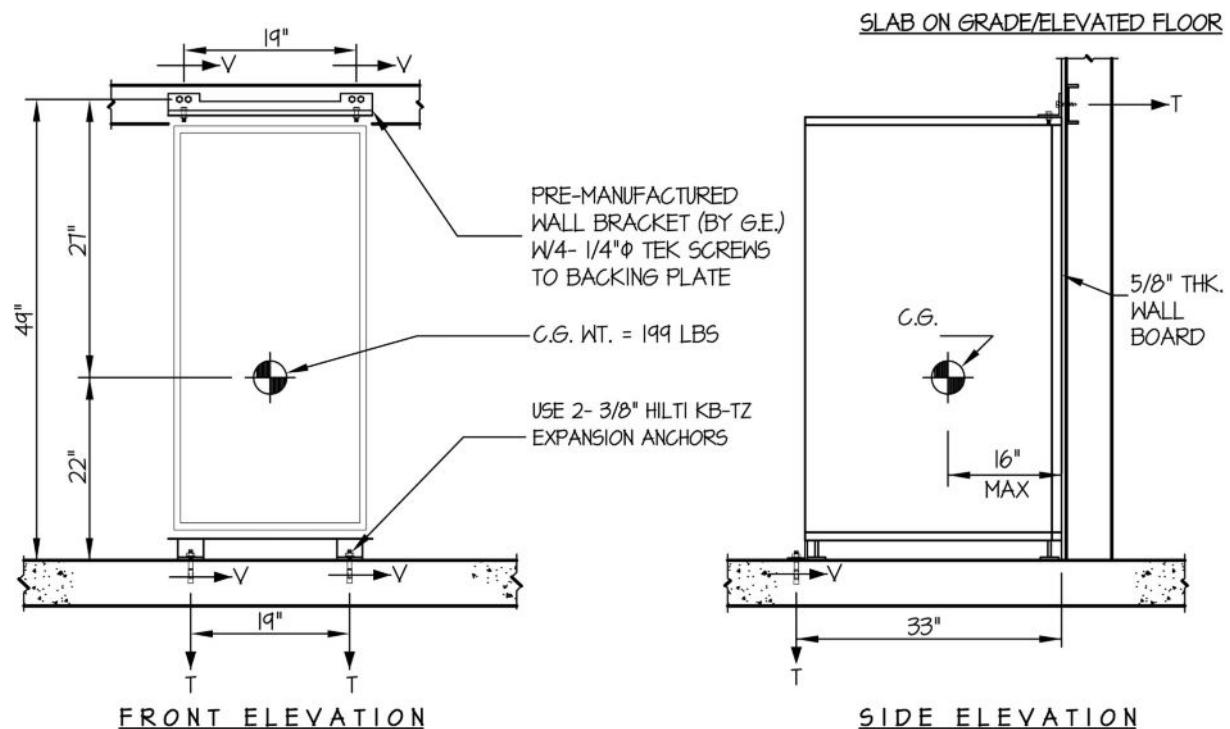


Illustration 2-103: Large Display Monitor suspension (LD option)

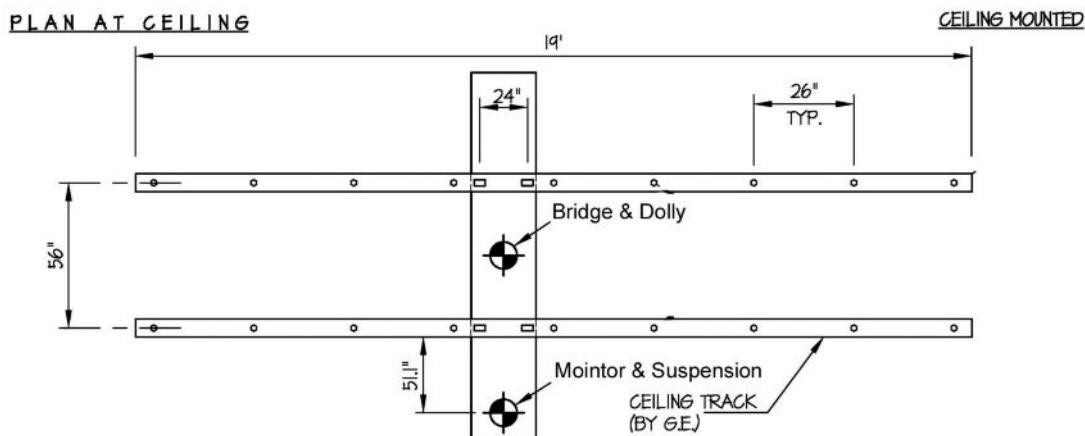
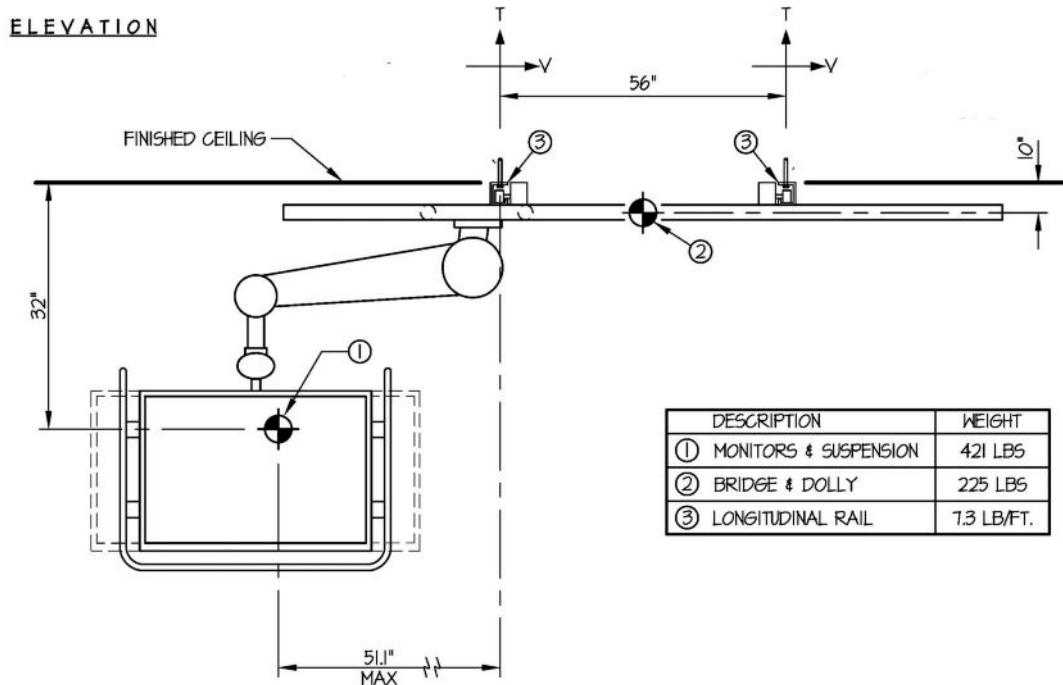
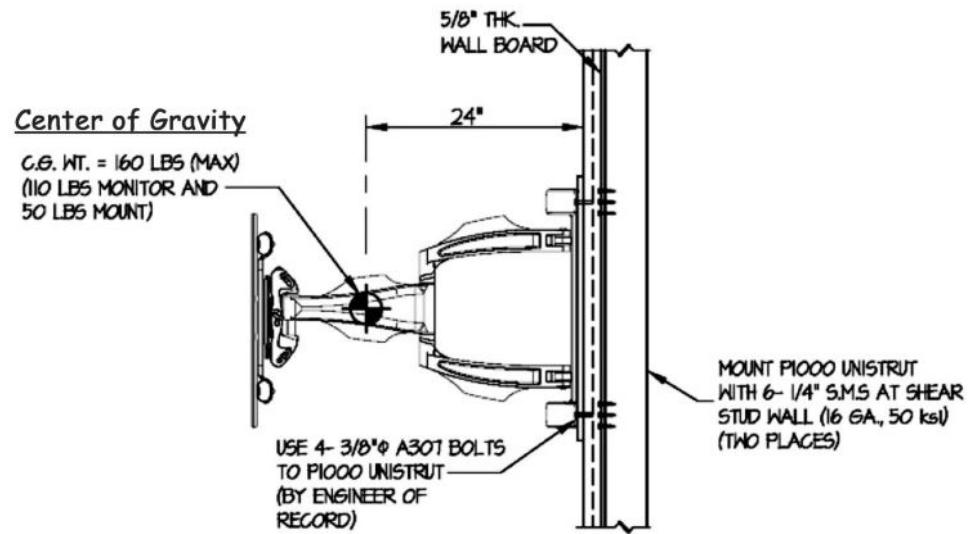


Illustration 2-104: Large Display secondary monitor Swing out arm (LD option)



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Chapter 3 Special Construction Requirements

1 Radiation Protection

Because x-ray equipment produces radiation, special precautions may be needed or special site modifications may be required. The General Electric Company does not make recommendations regarding radiation protection. It is the purchasers' responsibility to consult a radiation physicist for advise on radiation protection in x-ray rooms.

2 EMI Consideration

IEC60601-1-2 Electromagnetic Standard Compliance & Documentation

The information contained in this section is also found in the Innova system Operator Manual.

2.1 General Scope

This equipment complies with IEC-60601-1-2: Edition 2.1 and Edition 3 standard for medical devices

The Innova system is suitable to be used in the electromagnetic environment, as per the limits & recommendations described in the tables here after:

- Emission Compliance level & limits ([Table 3-1](#)).
- Immunity Compliance level & recommendations to maintain equipment clinical utility (see [Table 3-2](#),[Table 3-3](#) and [Table 3-4](#)).

NOTE: This system complies with above-mentioned EMC standard when used with supplied cables up to maximum lengths referenced in the MIS MAPS or system cables interconnect diagrams.

2.2 Electromagnetic Emission

The Innova system is intended for use in the electromagnetic environment specified below.

The customer or the user of the Innova system should assure that it is used in such an environment.

Table 3-1:

Emissions	Test Compliance	Electromagnetic Environment
Radio-Frequency Emissions CISPR11	Group1 Class A limits	The Innova system uses Radio Frequency energy only for its internal function. Therefore, its Radio Frequency emissions are very low and are not likely to cause any interference in nearby electronic equipment.
		The Innova system is suitable for use in all establishments other than domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Not applicable	The Innova system is suitable for use in all establishments other than domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Voltage fluctuations / flicker emissions IEC 61000-3-3	Not applicable	The Innova system is suitable for use in all establishments other than domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

2.3 Electromagnetic Immunity

2.3.1 Electromagnetic Immunity IEC 60601-1-2

The Innova system is intended for use in the electromagnetic environment specified below.

The customer or the user of the Innova system should assure that it is used in such an environment.

Table 3-2:

Immunity Test	IEC 60601-1-2 Test Level	Compliance Level	Electromagnetic Environment
Electrostatic discharge (ESD) IEC 61000-4-2	+/-6 kV contact +/-8 kV air	+/-6 kV contact +/-8 kV air	Floors are wood, concrete or ceramic tile or floors are covered with synthetic material and the relative humidity is at least 30 %.
Electrical fast transient/burst IEC 61000-4-4	+/-2 kV for power supply lines +/-1 kV for input/output lines	+/-2 kV for power supply lines +/-1 kV for input/output lines	Mains power quality is that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	+/-1 kV line(s) to lines(s) +/-2 kV line(s) to earth	+/-1 kV line(s) to lines(s) +/-2 kV line(s) to earth	Mains power quality is that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0 % U_n for 5 sec	0 % U_n for 5 sec	Mains power quality is that of a typical commercial or hospital environment. If the user of the Innova system requires continued operation during power mains interruptions, it is recommended that the Innova system be powered from an uninterruptible power supply.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields is at levels characteristic of a typical location in a typical commercial or hospital environment.
Note: U_n is the AC mains voltage prior to application of the test level.			

The Innova system is intended for use in the electromagnetic environment specified below.

The customer or the user of the Innova system should assure that it is used in such an environment.

Table 3-3:

Immunity Test	IEC 60601-1-2 Test Level	Compliance Level	Electromagnetic Environment
Conducted Radio Frequency IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	$V_1 = 3 \text{ V}$	Portable and mobile RF communications equipment is used no closer to any part of the Innova system, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
Radiated Radio Frequency IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz See statement below for Large Permanently Installed Medical Equipment.	$E_1 = 3 \text{ V/m}$ See statement below for Large Permanently Installed Medical Equipment.	<p>Recommended separation distance: $d = [3.5/V_1]\sqrt{P}$ $d = [3.5/E_1]\sqrt{P}$, from 80 MHz to 800 MHz $d = [7/E_1]\sqrt{P}$, from 800 MHz to 2.5 GHz</p> <p>Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m).</p> <p>Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey*, are less than the compliance level in each frequency range**.</p> <p>Interference may occur in the vicinity of equipment marked with the following symbol:</p>  <p>with the following symbol:</p>

NOTE: * Field strengths from fixed transmitters, such as base stations for cellular telephones and land mobile radios, amateur radio, AM and FM radio broadcast, and TV broadcast cannot be estimated accurately. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be performed. If the measured field strength exceeds the RF compliance level above, observe the Innova system to verify normal operation in each use location. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the Innova system

** Over the frequency range 150 kHz to 80 MHz, field strengths are less than 3 V/m.



WARNING

THE INNOVA IGS SYSTEM IS A LARGE, PERMANENTLY-INSTALLED MEDICAL EQUIPMENT FOR WHICH THE SIMULATED OPERATION IN AN ANECHOIC CHAMBER IS NOT FEASIBLE AND CONSEQUENTLY IS EXEMPT FROM THE TESTING REQUIREMENT SPECIFIED BY IEC 61000-4-3.

The Innova IGS system has not been tested for radiated RF IMMUNITY over the entire frequency range 80 MHz to 2.5 GHz.

The Innova IGS system has been tested in situ for radiated RF IMMUNITY only at selected frequencies in the range 80 MHz to 2.5 GHz.

ISM Frequency (MHz)	Field Level	Modulation
433.920 (ISM) ⁽¹⁾	3 V/m	80 % AM at 1 kHz rate
915 (ISM) ⁽¹⁾		
1440		
1750		
1920		
2450 (ISM) ⁽¹⁾		

NOTE: ⁽¹⁾: Industrial, Scientific and Medical (ISM) radio bands.

Equipment used for tests:

- RF signal generator,
- RF power amplifier,
- Transmitting antenna,
- Field sensor,
- Field meter.

The Recommended Separation Distances are listed in [Table 3-4](#).

These are guidelines. Actual conditions may vary.

2.3.2 Recommended Separation Distances for Portable and Mobile RF Communications Equipment IEC 60601-1-2

Table 3-4:

Frequency of Transmitter	150 KHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz
Equation	$d = [3.5 / V_1] \sqrt{P}$	$d = [3.5 / E_1] \sqrt{P}$	$d = [7 / E_1] \sqrt{P}$
Rated Power of Transmitter (watts)	Distance (meters)	Distance (meters)	Distance (meters)
10 mW	0.11	0.11	0.22
100 mW	0.37	0.37	0.74
1	1.1	1.1	2.3 (*)
10	3.7	3.7	7.4
100	12	12	23

For transmitters rated at a power not listed above, the DISTANCE can be estimated using the equation in the corresponding column, where P is the power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE: These are guidelines. Actual conditions may vary.

2.4 Limitations Management

Adhering to the distance separation recommended in [Table 3-4](#), between 150 KHz & 2.5 GHz, will reduce disturbances recorded at the image level, but may not eliminate all disturbances.

However, when installed and operated as specified herein, the system will maintain its essential performance by continuing to acquire, display, and store diagnostic quality images safely.

For example, a 1W mobile phone (800 MHz to 2.5 GHz carrier frequency) shall be put 2.3 meters (see (*) [Table 3-4](#)) apart from the Innova system (in order to avoid images interferences risks).

2.5 Use Limitation



WARNING

THE USE OF ACCESSORIES, TRANSDUCERS, AND CABLES OTHER THAN THOSE SPECIFIED MAY RESULT IN DEGRADED ELECTROMAGNETIC COMPATIBILITY OF THE INNOVA SYSTEM.

In case of an Innova system, the Gantry (digital detector) shall be apart 1 meter from the generator cabinet, and 1 meter apart from the analog (CRT) monitors. These distances specifications shall minimize the low frequency magnetic field interference risk.

Other electrical equipment may disturb and interfere with these Innova components. The control of the clearing distances from the noise sources is recommended from the HF electrosurgery generator, power supplies converters from nearby monitors or from other close electrical equipment). Refer to respective device manufacturers instructions & recommendations in such cases.

2.6 Installations Requirements & Environment Control



NOTICE

In order to minimize interference risks, the following requirements shall apply:

- Cables shielding & grounding:

All interconnect cables to peripheral devices must be shielded and properly grounded. Use of cables not properly shielded and grounded may result in the equipment causing radio frequency interference.

- Separated Power supply distribution panel & separated power line:

- This product complies with the radiated emission limits as per the CISPR11 Group1 ClassA standard.
- The Innova system is predominantly intended for use (e.g. in hospitals) with a dedicated supply system, and with an X-ray shielded room.
- In case of using in a domestic environment (e.g. doctors' offices), in order to avoid interferences, it is recommended to use a separated AC power distribution panel & separated power line, and an X-ray shielded room.

- Subsystem & accessories Power supply distribution:

All components, accessories subsystems, systems which are electrically connected to the Innova system, have to be all AC power supplied by the same power distribution panel & line.

NOTE: In order to avoid interferences, the same AC power distribution panel should supply all components, accessories, the Innova system (& subsystems as the Advantage Workstation). The separated AC power line should supply the panel.

- Stacked components & equipment:

The Innova system should not be used adjacent to or stacked with other equipment; if adjacent or stacked use is necessary, the Innova system should be observed to verify normal operation in the configuration in which it will be used.

- Low frequency magnetic field:

In case of an Innova system, the Gantry (digital detector) shall be apart 1 meter from the X-Ray generator cabinet, 1 meter from the PDB cabinet, 3 meters from the UPS cabinet and 1 meter apart from monitors. These distances specifications shall minimize the low frequency magnetic field interference risk.

- Static Magnetic Field Limits : The Innova system is compatible with the Earth magnetic Field. Earth magnetic Field is lower than 1 Gauss.

- Electrostatic discharges environment & recommendations:

- In order to reduce electrostatic discharge interference, install a charge dissipative floor material to avoid electrostatic charge buildup.

- The relative humidity shall be at least 30 percent.
- The dissipative material shall be connected to the system ground reference, if applicable.



NOTICE

Route separately EMI filter incoming power lines and outgoing power lines (refer to *Electrical Connections*).

NOTE: The maximum distance between the EMI filter and the PDB is 3 m.

NOTE: The Fluoro UPS (CE) option requires an EMI filter box placed upstream the PDB. The EMI filter box will be delivered as a single product by GE Consumer & Industrial.

Chapter 4 Environmental Requirements

1 Relative Humidity and Temperature



NOTICE

Avoid extremes in temperatures

Innova system room climate requirements – relative humidity and temperature (individual products or components are classed by their installation area for **In use** see -[Table 4-1](#))

Table 4-1:

INSTALLATION ROOM OF PRODUCT OR COMPONENT	RELATIVE HUMIDITY (NON-CONDENSING)		TEMPERATURE			
	IN-USE		IN-USE (See Note (2a))		RECOMMENDED (See Note (2b))	
	MIN	MAX	MIN	MAX	MIN	MAX
Examination room	30%	70%	+15°C +59°F	+32°C +90°F	Design for Patient/ Operator Comfort	
Technical room (See Note (1))	30%	75%	+10°C +50°F	+30°C +86°F	+13°C +55°F	+25°C +77°F
Technical room with Fluoro UPS optional	30%	75%	+20°C +68 °F	+25°C +77°F	+20°C +68°F	+25°C +77°F
Control room	30%	75%	+15°C +59°F	+35°C +95°F	+20°C +68°F	+25°C +77°F

NOTE: (1): The target temperature (best recommended) is 18°C (64°F).

NOTE: (2a): **In use temperature limits** specify the range where the system shall work. Operating outside these limits could occur severe performance and reliability issues.

(2b): **Recommended temperature limits** specify the range where it is recommended to adjust air conditioning control in order to warranty current operations inside the in use range.

Relative Humidity and Temperature: Refer to [Table 4-1](#). To obtain relative humidity and temperature requirements for components not specified in the table, refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova System Compatibility](#).



NOTICE

In some cases condensation occurs and water drops from outlets and pipes of the air conditioner in the technical room.

Therefore, it is critical to install the cabinets where there is no risk of flood from the air conditioner.



NOTICE

Ensure the air outlet positions in the exam room is not in area closed to monitor suspension carriage/rail.

IVUS Option :

1. IVUS Rev 2 and Rev 2+

Table 4-2:

Humidity (non condensing) :	30 - 80 %	
Temperature (at 20% Relative Humidity) :	5 - 35 C	(41 - 95 F)

2. IVUS Rev 3

Refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual.](#)

2 Altitude and Atmospheric Pressure

Refer to [Table 4-3](#). To obtain altitude and atmospheric pressure requirements for components not specified in [Table 4-3](#), refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova System Compatibility](#).

Table 4-3: Altitude and Atmospheric Pressure

INSTALLATION ROOM OF PRODUCT OR COMPONENT	ALTITUDE (meters)		ATMOSPHERIC PRESSURE (kPa)	
	IN-USE		IN-USE	
	MIN	MAX	MIN	MAX
Examination room	0	2000	79.4	106
Technical room (See Notice below)	0	2000	79.4	106
Control room	0	2000	N/A	N/A



NOTICE

The chiller is able to dissipate maximum continuously power at 3000 meters in an ambient temperature up to 20°C, for this altitude the technical room temperature shall not exceed 20°C.

3 Heat Output

3.1 Equipment Heat Output tables

Refer to [Table 4-4](#). To obtain heat output information for components not specified in [Table 4-4](#), refer to the appropriate component Pre-Installation Manual listed in [Chapter 2, Basic Innova System Compatibility](#).

Table 4-4:

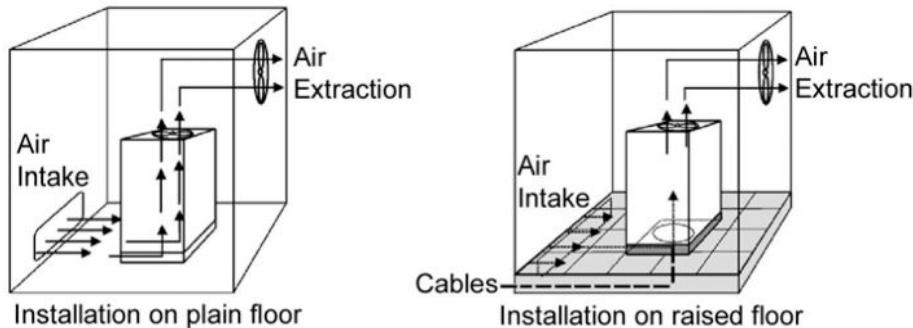
		HEAT OUTPUT							
		Stand by		Moderate Use (4)		Typical Use (4)		Maximum Use (4)	
Room	Core System	kW	BTU/hr	kW	BTU/hr	kW	BTU/hr	kW	BTU/hr
Exam Room	Innova LC positioner and table	0.41	1394	0.55	1858	0.89	3020	1.62	5517
Ctrl Room	DL user area with 1 TFT monitor	0.16	546	0.16	546	0.16	546	0.16	546
	2 B&W flat monitors	0.17	573	0.17	573	0.17	573	0.17	573
Tech Room	C1 Cabinet	0.41	1398	0.69	2366	0.99	3389	1.29	4412
	C2 Cabinet	0.29	989	0.33	1125	0.54	1828	0.87	2966
	Coolix X-Ray tube chiller (1) (2)	2.53	8619	4.49	15309	5.49	18725	6.93	23625
	Detector conditioner	0.21	709	0.21	709	0.21	709	0.21	709
	Main disconnect panel - PDB	0.4	1534	0.45	1534	0.45	1534	0.45	1534
Total for core system		4.6	15762	7.0	24020	8.9	30324	11.7	39881
<hr/>									
Room	Options (3 & 5)	Stand by		Moderate Use (4)		Typical Use (4)		Maximum Use (4)	
Exam Room	3 in room B&W TFT monitors	0.25	859	Same values as Stand by		Same values as Stand by		Same values as Stand by	
	In room AW TFT monitor	0.12	409						
	Typical injector	0.09	320						
	LD Monitor	0.5	1706						
Ctrl Room	AW work station	0.35	1201	Same values as Stand by		Same values as Stand by		Same values as Stand by	
	2 AW TFT monitors	0.24	818						
	Printer	0.31	1054						
	LD Monitor (6)	0.5	1706						
Tech Room	LD Cabinet	0.15	512						
	3kVA Cabinet UPS - model 9130 (7)	0.37	1257						
Typical configuration without fluoro UPS		5.9	20424	8.4	28682	10.3	34985	13.1	44543

NOTE: (1) Air flow requirements 1200 m³/h (706 CFM)

- NOTE: (2) For more details, consult appropriate pre-installation manual
- NOTE: (3) For UPS 20 kVA option refer to [Section 3.2](#)
- NOTE: (4) **Moderate use:** 8 cases / 10 hours, **typical use:** 11 cases / 10 hours, **maximum use:** during case.
- NOTE: (5) For IVUS environmental limitations and heat output, refer to [Section 3.3](#)
- NOTE: (6) The 2nd optional LD monitor is not necessarily in the control room. It may be installed in the exam room (outside patient vicinity).
- NOTE: (7): On batteries heat output = 0.52 kW / 1755 BTU

3.2 Fluoro UPS Option

Illustration 4-1:



NOTICE

The Fluoro UPS batteries require a cooling system to keep ambient temperature below 25 °C (77 °F).

The heat produced by the UPS is transferred to the environment by its ventilation. Cooling air enters the cabinets through the air inlet (grids) located at the bottom and exhausted through the outlet on the roof. A suitable ventilation or cooling system must be installed to extract the heat from the UPS room.



CAUTION

Make sure there is a ventilation air flow, preferably ensured by natural air flow, otherwise by enforced ventilation, so that hydrogen concentration is below 1% (according to Standard IEC 62040-1-2).



NOTICE

Do not put anything on the top of the cabinet.

If the UPS is placed on a raised floor, the airflow for UPS cooling should enter from underneath the UPS, through the appropriate aperture on the raised floor.

If the UPS runs in a dusty environment, we recommend strongly to install filters on the air inlet of the UPS room. In this case it should be considered that these filters can cause reduced speed at the air inlet.

The size of the air inlet has therefore to be dimensioned accordingly.

Contact your Local Distributor or one of the Service Centre, which will help you to find valuable solutions.

The tables below indicate the heat dissipation at full load at **PF = 0.8 lag**, and charged battery, up to 1000 m (3280 ft) altitude, for cooling air 25°C (77°F) to 30°C (86°F).

- **VFI** (Voltage Frequency Independent) UPS system where the load is continuously supplied by the inverter through the rectifier.
- **SEM** (Super Eco Mode) permitting the maximum energy saving.

Table 4-5: Fluoro UPS CE

Losses		Cooling air flow	
VFI	SEM	VFI	SEM
2.14 kW	0.64 kW	625 m³/h	190 m³/h

Table 4-6: Fluoro UPS UL

Losses		Cooling air flow	
BTU / hr	kW	CFM	m³/h
6751	1.98	301	512



NOTICE

If installed in the technical room, the UPS may impact on its layout. It may also be installed in a separate room. This depends on hospital constraints, local regulations or EHS rules. Clearance, weight of UPS, airflow and cooling system should be adapted for the UPS.



WARNING

ELECTRICAL CONTRACTOR IS RESPONSIBLE FOR PROVIDING AND CONNECTING THE CABLES FROM THE PDB TO THE UPS AND CONFIGURING THE PDB IN BYPASS MODE. GEHC IS RESPONSIBLE FOR POWERING ON THE SYSTEM WITH THE UPS IN BYPASS MODE. GEDE IS RESPONSIBLE FOR UPS COMMISSIONING.



CAUTION

The Fluoro UPS can be installed either in an Electrical Local with restricted access or in the innova technical room as per local regulations.



NOTICE

Make sure that local regulations have been applied for the installation of the Fluoro UPS (dedicated room/fire detection etc.)

NOTE: Refer to the UPS vendor Service Operating Manual for more details:

- [UL Model: UPS Option - Operating Manual SG-UL Series - 20 kVA](#)
- [CE Model: UPS Option - Operating Manual SitePro 20 kVA \(series 8\)](#)

3.3 IVUS Option

Device	Location	Heat output
PC	Control room	335W – 1206 BTU/h
Monitor	Control room	125W – 427 BTU/h
Printer	Control room	67W – 229 BTU/h (idle)

NOTE: For IVUS Rev 3, refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).



WARNING

TAKE ANY STEPS NECESSARY TO PROVIDE CORRECT TEMPERATURE AND DUST FREE ENVIRONMENT FOR CORRECT OPERATION OF PC IN THE CONTROL ROOM.

3.4 Large Display Subsystem Option



NOTICE

The Large Display Cabinet requires a cooling system to keep ambient temperature below 35 °C (95 °F).

The heat produced by the Large Display Cabinet is transferred to the environment by its ventilation. Cooling air enters the Large Display cabinet through the air inlet (grids) located at the front and exhausted through the outlet on the rear & from underneath the Large Display Cabinet. A suitable ventilation or cooling system must be installed to extract the heat from the Large Display Cabinet room

Ventilation holes must not be covered or closed; otherwise the heat generated in the device cannot be dissipated sufficiently.

Avoid dusty environments:

The Large Display Monitor has been designed for use in the clean environment of medical diagnostics. The Large Display Monitor dissipates heat through the openings at the rear. Dust from dirty environments can penetrate into the display through these openings. In the extreme case, deposits are possible which become evident as dark spots in a white picture and which can result in deterioration of the luminance. Protect the display from dust, e.g. during building measures at the installation location, and use the original packaging for transport.



NOTICE

Do not put anything on the top of the Large Display cabinet.

If the Large Display Cabinet runs in a dusty environment, we recommend strongly installing filters on the air inlet of the Large Display Cabinet room. In this case it should be considered that

these filters can cause reduced speed at the air inlet. The size of the air inlet has therefore to be dimensioned accordingly.



NOTICE

If installed in the technical room, the Large Display Cabinet may impact its layout. It may also be installed in a separate technical room. This depends on hospital constraints, local regulations or EHS rules. Clearance, weight of Large Display Cabinet, airflow and cooling system should be adapted for the Large Display Cabinet.

Make sure that local regulations have been applied for the installation of the Large Display Cabinet (dedicated room/fire detection etc.)

NOTE: Refer to the Large Display Subsystem vendor Service manual for more details



NOTICE

Ventilation instructions:

- The Large Display Cabinet must be placed in a sufficiently ventilated area; the ambient temperature should not exceed 95°F (35°C).
- Clearance around the front of the unit should be sufficient to enable free passage of personnel with the doors fully open, and to allow sufficient airflow to the door vents – Min. 100cm / 40”
- It is important that air can move freely around and through the LD Cabinet. Do not block the air vents.
- Avoid locations in direct sunlight or near heat sources.

3.5 Vivid E9 Ultrasound Option

Heat Dissipation for Vivid E9 system is 3800 BTU/h.

4 Acoustic Specifications

- Less than 50 dB (A) at 1 meter for Gantry.
- Limited to 50 dB (A) at 1 meter for Omega IV, Omega V tables.
- Limited to 58 dB (A) at 1 meter for Tilting table.
- Limited to 55 dB (A) at 1 meter for C2 Cabinet.
- Limited to 60 dB (A) at 1 meter for the COOLIX 4100.
- Limited to 65 dB (A) at 1 meter for C1 Cabinet.
- Limited to 52 dB (A) (background of 35 dB (A)) at 1 meter for Digital Detector Chiller Thermo-Con.
- Less than 50 dB (A) at 1 meter for a DL LCD monitor.
- Less than 60 dB (A) at 1 meter for the Fluoro UPS.
- less than 55 dB(A) at 20 degrees Celsius, measured in the operators head position, 20 cm in front of the keyboard's right corner, at 1.30 m above the floor, and in a distance of 1 meter at all four sides, 1 meter above the floor for the Vivid E9 ultrasound unit.

NOTE: Both cabinets C1 and C2 generate 70 dB noise altogether. Noise can be reduced if cabinets are slightly separated, and as far as possible from Exam room.

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Chapter 5 Electrical Requirements

1 Power Requirements

1.1 System Electrical Characteristics

NOTICE

It is the customer's responsibility to ensure that electrical installation is compliant to local regulations. To avoid risk of electric shock, this equipment must only be connected to a supply mains with Protective Earth.

Power supply and ground cables shall be dedicated to the system. They must not be used to supply other systems.

Power supply and ground cables shall be kept separated from room system cables.

Power supply and ground cables must be connected to the same distribution panel. They must run near one to the other.

Power supply and ground cables provided by the customer shall be compliant with local regulations (e.g. UL, NFPA 70, CSA, IEC, CCC).

1.1.1 Core system

Table 5-1:

Nominal voltage	Frequency	Power consumption		Type of power input	
		Nominal	Peak	Without Fluoro UPS option	With Fluoro UPS option
380 V ± 10%					
400 V ± 10%					
415 V ± 10%					
480V ± 10%	60 Hz only (± 3 Hz)	60 kVA	150 kVA	3~	3N~

The Hospital circuit breaker should fit the current protection of the Innova system:

- 150A/ 480V, 3 Phases for UL
- 80A/ 380V, 3 Phases for CE



WARNING

THIS FLUORO UPS IS ONLY DESIGNED TO OPERATE IN A WYE-CONFIGURED ELECTRICAL SYSTEM WITH A SOLIDLY GROUNDED NEUTRAL.

FOR MORE DETAILS, REFER TO THE FLUORO UPS DOCUMENTATION.

1.1.2 Options

Table 5-2:

Option	Nominal voltage	Frequency	Nominal Power consumption	Type of power input
LDM	100-120 V / 220-240 V	50 Hz or 60 Hz (± 3 Hz)	3 kVA	Single phase
AW	100-127 V / 200-240 V	50 Hz or 60 Hz	11 A / 5.5 A	Single phase
S5I GE	100-120 V / 230 V	50 Hz or 60 Hz	400 VA	Single phase



! WARNING

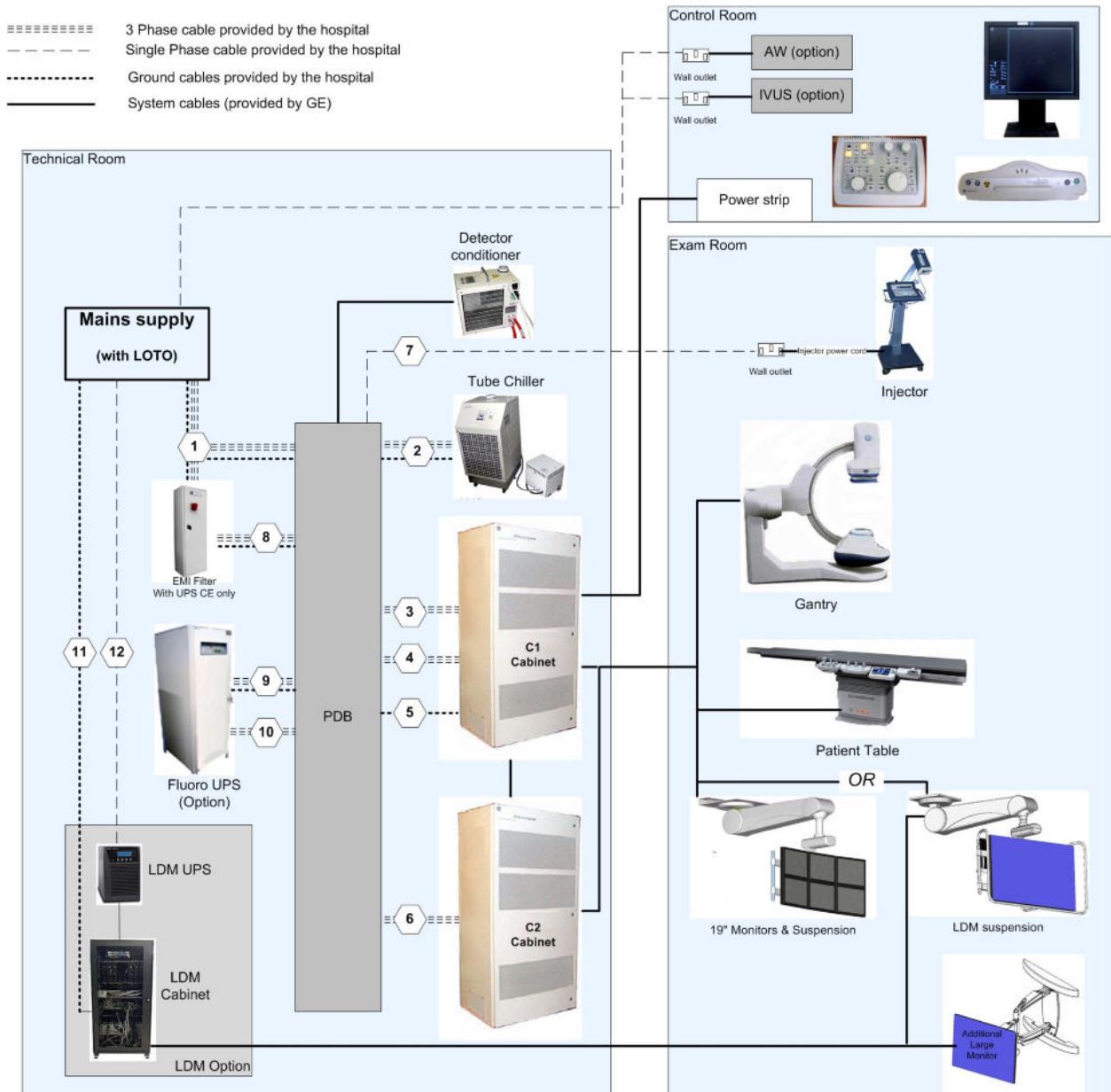
PRIOR TO EACH INSTALLATION, ENSURE THAT THE ECG POWER CABLE IS CONNECTED TO A LINE THAT IS PROTECTED AGAINST SHORT CIRCUIT HAZARDS, ACCORDING TO LOCAL REGULATIONS AND THAT THE ECG MONITOR IS POWERED ACCORDING TO IEC60601-1 REQUIREMENTS (LEAKAGE CURRENT AND GROUNDING).

1.2 Power and grounding

The power cords between the peripherals in the control room and the power strip are not shown on the following diagram. All are single phase power cords (with phase, neutral line and protective Earth) and are provided by GE.

1.2.1 Schematic

Illustration 5-1: Electrical and grounding schematic



NOTE: For the complete system connection schematic, refer to the MIS Map and MIS Chart provided in the Advanced Service Manual.

1.2.2 Cables to be provided by the installer

For cable # in the following tables, refer to [Illustration 5-1](#)

1.2.2.1 Core system

Table 5-3: Core system cables

Cable #	From	To	Type	Characteristics	Comments
1	Mains	PDB	3 phases	Diameter in conformity with the Max Line Impedance table	
			Neutral	Diameter in conformity with the Max Line Impedance table	
			Ground	AWG2/35 mm ² , and not smaller than the neutral conductor	
2	PDB	Tube Chiller auto transformer	3 phases	Max length 24 m between PDB and chiller.	
			Ground	Min diameter AWG12 (UL) / 4 mm ² (CE)	
3	PDB	C1 cabinet	3 phases	Max length 12m. Min diameter AWG10 (UL) / 4 mm ² (CE)	
4	PDB	C1 cabinet	3 phases	Max length 12 m. Diameter in conformity with the Max Line Impedance below	For the X-ray generator inside C1
5	PDB	C1 cabinet	Ground	Max length 12 m. Min diameter AWG2/35 mm ² .	
6	PDB	C2 cabinet	3 phases	Max length 12 m. Min diameter AWG10	Only for UL (system cable from C1 to C2 provided for CE)
7	PDB	Injector wall outlet	Phase, neutral and ground	Max length 24 m. Min diameter AWG12 (UL) / 1.5 mm ² (CE)	120V outlet UL

Table 5-4:

Max Line Impedance for the line from the Hospital outlet to the X-rays Generator in C1 cabinet (cables #1 and #4)						
V	380	400	415	440	460	480
Ω	0.09	0.096	0.102	0.108	0.114	0.12

1.2.2.2 For the Fluoro UPS option

Table 5-5: Fluoro UPS option cables

Cable #	From	To	Type	Characteristics	Comments
1	Mains	PDB	3 phases	Diameter in conformity with the Max Line Impedance table	Only with UL Fluoro UPS option
			Neutral	Diameter in conformity with the Max Line Impedance table	
			Ground	AWG2/35 mm ² , and not smaller than the neutral conductor	
1	Mains	EMI Filter	3 phases	Diameter 35 mm ² min, in conformity with the Max Line Impedance table	Only with CE Fluoro UPS option
			Neutral	Diameter 35 mm ² min, in conformity with the Max Line Impedance table (35 mm ² min)	
			Ground	AWG2/35 mm ² and not smaller than the neutral conductor	
8	EMI Filter	PDB	3 phases	Max length 3 m. Diameter in conformity with the Max Line Impedance table	Only with CE Fluoro UPS
			Neutral	Max length 3 m. Diameter in conformity with the Max Line Impedance table	
			Ground	AWG2/35 mm ² , and not smaller than the neutral conductor	
9	PDB	Fluoro UPS	3 phases	10 mm ² (CE) AWG6 (UL)	Max diameter AWG3
				10 mm ² (CE) AWG6 (UL)	
			Neutral	10 mm ² (CE) AWG6 (UL)	
				10 mm ² (CE) AWG6 (UL)	
			Ground	10 mm ² (CE) AWG6 (UL)	
10	Fluoro UPS	PDB	3 phases	10 mm ² (CE)	Max diameter AWG3

NOTE: Cables 1 and 8 should be routed separately (CE Only) in order to avoid EMC interference.

1.2.2.3 For the LDM option

Table 5-6: LDM option cables

Cable #	From	To	Type	Characteristics	Comments
11	Mains	LDM cabinet	Ground only	UL: AWG10 (5.26 mm ²)	
				CE: AWG12 (2.5 mm ²)	

Cable #	From	To	Type	Characteristics	Comments
12	Mains	LDM UPS	Phase, neutral and ground	UL: AWG10 (5.26 mm ²)	
				CE: AWG12 (2.5 mm ²)	

1.2.3 LOTO devices provided by the installer

1.2.3.1 Core system

A wall circuit breaker or equivalent device with LOTO capability must be installed on the mains line to the PBD. This device must be compatible with the power input specifications of the system. The customer is responsible for the procurement, delivery and installation of this breaker.

1.2.3.2 LDM option

A wall circuit breaker or equivalent device with LOTO capability must be installed on the mains line to the LDM UPS. The rating of this device shall be 30A for UL and 16A for CE configurations. The customer is responsible for the procurement, delivery and installation of this breaker.

1.2.4 System interconnections

1.2.4.1 Emergency power off

The PDB (in technical room) is provided with an EPO button on its front panel. The customer is given the possibility to install additional EPO buttons (for instance in the exam room and in the technical room). The PDB is designed for 2 additional EPO buttons; these buttons shall be "Push to activate - Push to release" type, 2 contacts Normally Closed, compatible with 24 VDC.

The UL PDB is provided with 2 additional EPO buttons, the CE PDB is provided without additional EPO buttons.

The customer is responsible for the procurement, delivery and installation of the cables for these additional EPO buttons. The max length shall be 24 m, the recommended diameter is AWG14/2 mm².

Protect the Emergency Stop from accidental actuation.

1.2.4.2 X-Ray ON lights



NOTICE

THE X-RAY ON LAMP MUST BE INSTALLED IN THE EXAM ROOM IN CONFORMITY TO THE STANDARD IEC/EN 60601-2-43. THE X-RAY ON LAMP SHALL BE VISIBLE BY THE OPERATOR IN ALL THE LOCATIONS DEFINED FOR THE PERSONNEL WHO MAY RECEIVE SCATTERED RADIATION.

The system is designed to provide power for X-ray ON lights. The customer is responsible for the procurement, delivery and installation of the cables and the X-ray ON lights.

UL system:

The terminals 2 & 3 from the Positioner Bulkhead of the C2 cabinet shall be connected to PDB TB1 terminals 24 & 25 (cable provided by the installer, maximum length 24 m, minimum diameter AWG16). The lamp shall be 120V ~ 1A max, it shall be connected to PDB TB1 terminals 25 & 26, with cable AWG16 minimum

CE system:

The terminals 2 & 3 from the Positioner Bulkhead of the C2 cabinet shall be connected to PDB terminals BNC 12 & 13 (cable provided by the installer, maximum length 24m, minimum diameter 1.5 mm²). 2 slots are provided for 24V~ lamps or relays (PDB terminals 16-17 & 18-19), they shall be connected with cables of diameter 1.5 mm² minimum.

1.2.4.3 Room door interlock

The system is designed to provide room door interlocks that prevent X-ray emission when the door is open. IEC 60601-2-43 requires not to install door interlocks. It is the responsibility of the installer to check that this requirement is not in contradiction with local regulation. In case of conflict, the local regulation shall prevail.

To disable the door interlock: The terminals 8 & 9 from the Positioner Bulkhead of the C2 cabinet shall be shorted

To enable the door interlock: The terminals 8 & 9 from the Positioner Bulkhead of the C2 cabinet shall be connected to a door interlock switch provided by the installer. This switch shall be closed when the door is closed, it shall be compatible with 24 VDC. The max length between this switch and the C2 cabinet shall be 24 m

1.2.4.4 System on light

The system provides power for a System ON light. The customer is responsible for the procurement, delivery and installation of the cables and the X-ray ON lights.

For CE systems, the lamp shall be 230V~, 2 A max, the cables diameter shall be minimum 0.75 mm².

For UL systems, the lamp shall be 120V~, 250mA max, the cables diameter shall be minimum AWG18

The maximum length shall be 36 m

1.2.4.5 Injector

The PDB provides a terminal block that can power an injector. The customer is responsible for the procurement, delivery and installation of the cables from the PDB and of the wall outlet to power the injector. The maximum length of these cables shall be 24 m.

For CE systems, the PDB output is 230V~, 6 A max. The cables diameter shall be minimum 0.75 mm².

For UL systems, the PDB output is 120V~, 16A max, the cables diameter shall be minimum AWG12.

The impedance of the ground cable between the outlet and the PDB shall be 0.4 ohms maximum.

This wall outlet shall comply with the following requirements:

- It shall comply with the requirements of CSA C22.2 No. 42 & CSA C22.2 No. 49 (UL) or IEC 60884-1 (CE).
- Its rating shall be compatible with the max output of the PDB.
- It shall be of class I construction and the protective earth conductor shall be connected to the Earthing contacts in the socket-outlets.

- The creepage distances and air clearances shall comply with the following table:

Table 5-7:

System type	Between phase and Neutral		Between phase and ground and between neutral and ground		Between phase, Neutral and external accessible parts	
	Creepage distance	Air clearance	Creepage distance	Air clearance	Creepage distance	Air clearance
UL	2 mm	1 mm	1.5 mm	2.1 mm	3 mm	4.2 mm
CE	3 mm	1.6 mm	2.5 mm	2.2 mm	5 mm	4.4 mm



NOTICE

If the wall outlet is installed in the exam room, installation shall be done as per local regulation.

Additional multiple socket-outlet or extension cord shall not be connected to the terminal block or to the outlet. This outlet shall only power an injector; connecting other equipment could lead to fire hazard or excessive leakage currents.

The outlet shall be marked with the safety sign [Illustration 5-2](#) such that it is visible in normal use , with the maximum allowed continuous output in amperes or volt-amperes, and with the name of the equipment that will be connected (e.g. "injector".)

Illustration 5-2: Safety sign



1.3 Other Options

1.3.1 For USA only

A purchasable option I-sense (catalog number E4504B) allows the monitoring of the hospital main power line. It is recommended to install this option everywhere RMS and waveform variation events can impact the standard behavior of the system. I-sense is connected to each phase conductor and the ground. An analog telephone line also needs to be line to I-sense.

1.3.2 Third party monitors on suspension



WARNING

THIRD PARTY MONITORS INSTALLED ON THE SUSPENSION MUST NOT BE POWERED BY THE SYSTEM. DANGEROUS VOLTAGE MAY BE PRESENT AT THE INPUT OF THESE MONITORS, EVEN WHEN PDB IS OFF



NOTICE

THIRD PARTY MONITORS SHALL BE CERTIFIED IEC60601-1 OR IEC60950.

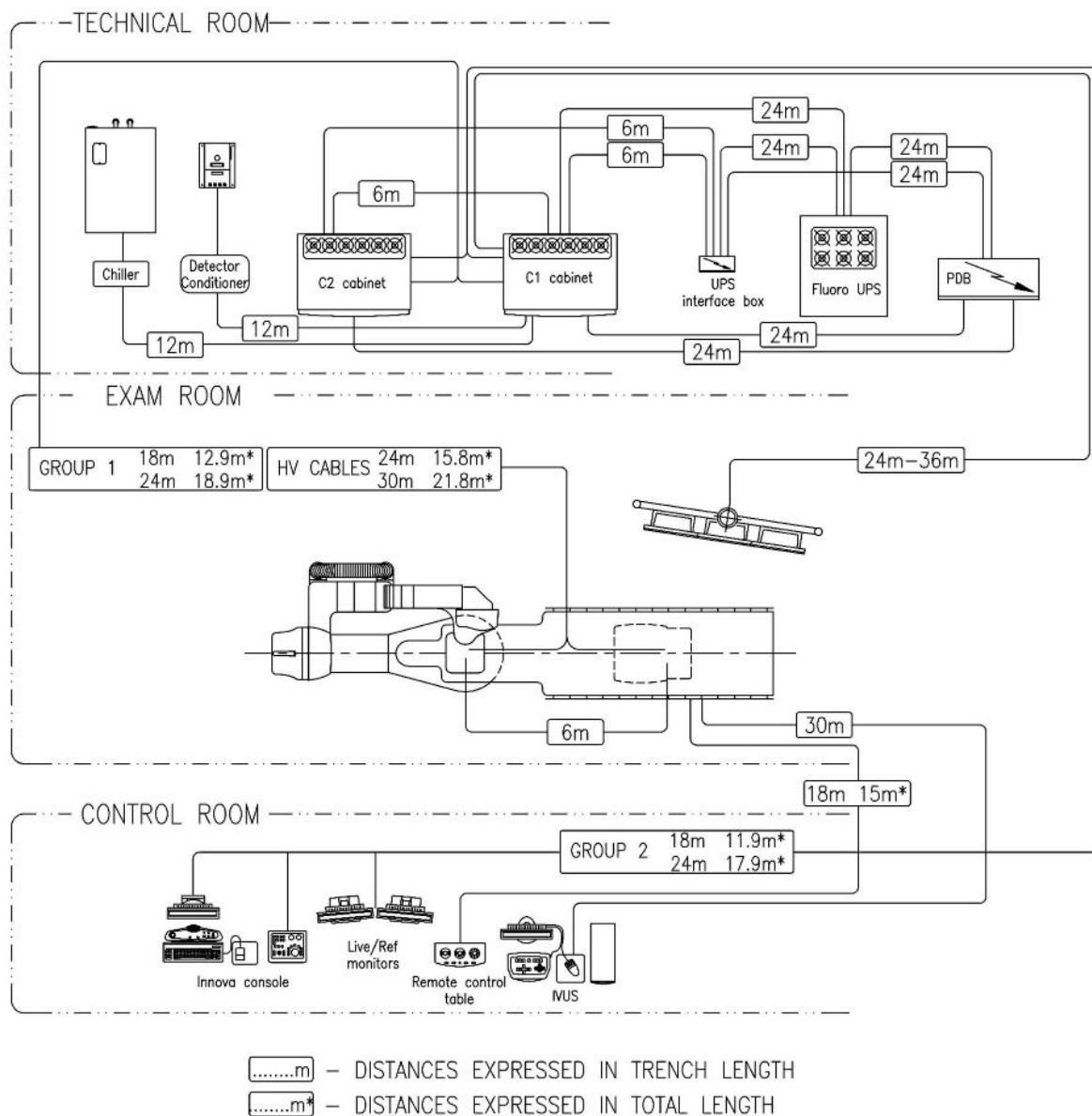
THIRD PARTY MONITORS MUST BE GROUNDED TO C1 CABINET GROUND BAR BY AN ADDITIONAL GROUND WIRE, AWG10 MINIMUM, AND 600V OF ANY UL TYPE.

2 System Cable Information

2.1 Physical Runs

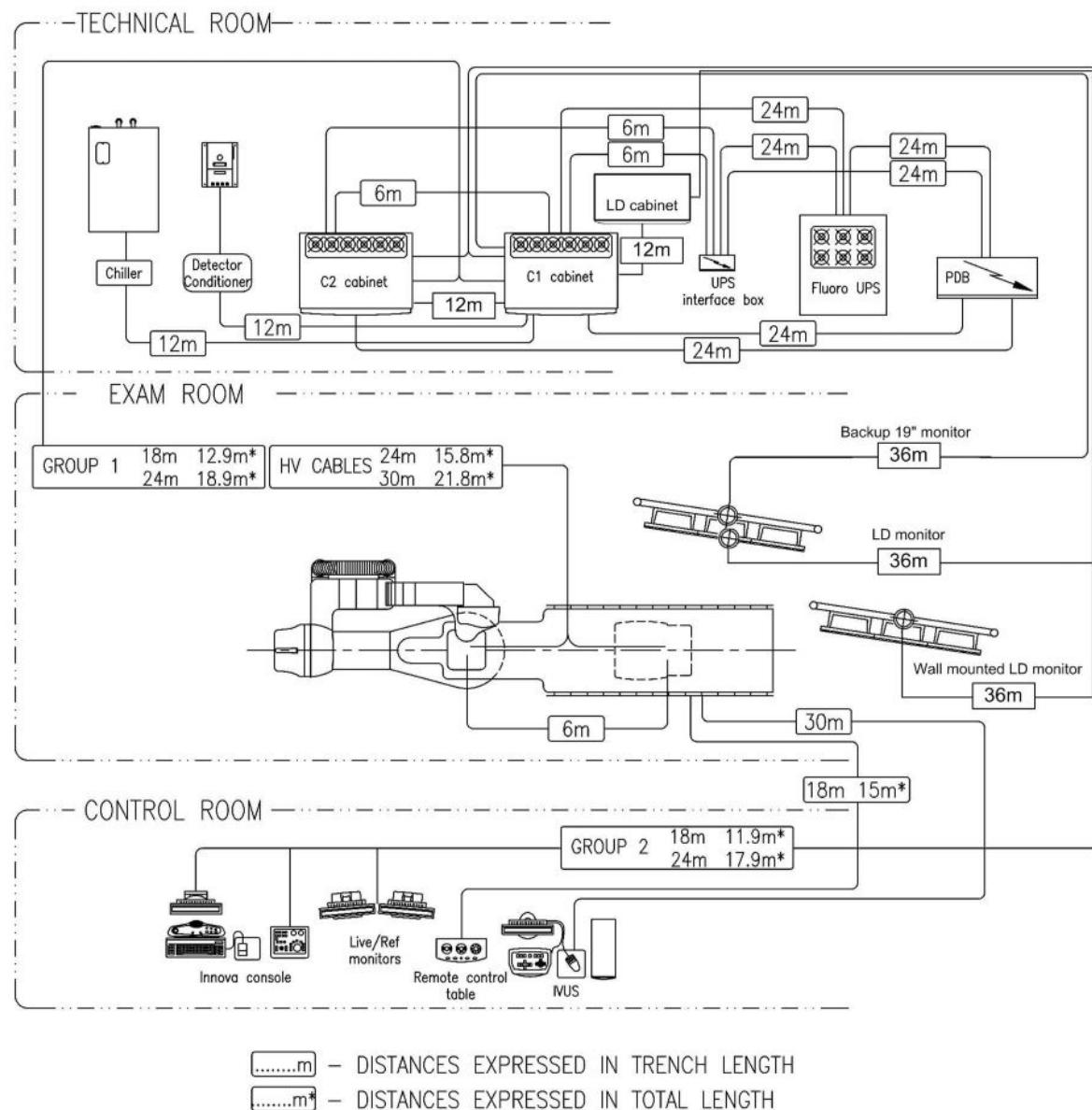
2.1.1 Physical Run Synoptic

Illustration 5-3: Mini / Maxi Interconnection Length (system with LCD monitors)



NOTE: A 24 m Emergency UPS stop cable 27561A is available in FRU.

Illustration 5-4: Mini / Maxi Interconnection Length (system with Large Display Monitor option)



NOTICE

Radius of curvature of the cables that connect the 8MP Monitor: min 35 mm.

2.1.2 System Core Matrix



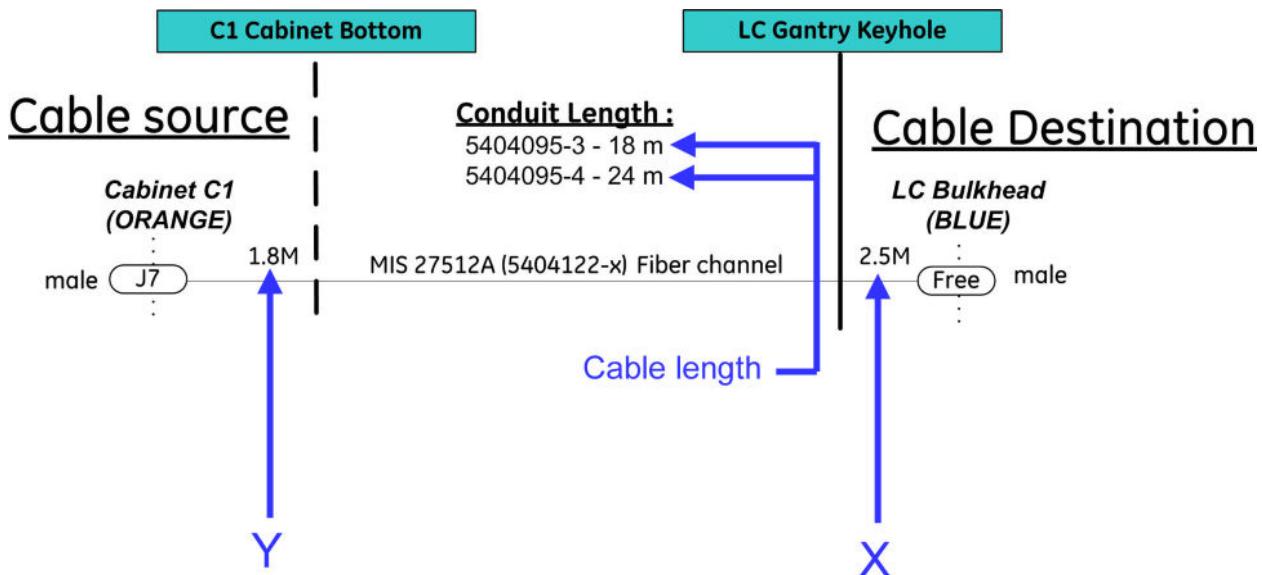
NOTICE

All lengths of cable are:

- in useable meter when you look at group level, or
- in meters (connector to connector) when you look at the cable level.

For a description of how to use the following cable group schematics, see below:

Illustration 5-5: Example of cable group schematic



Cable length data is as follows:

- **Cable Length** = the total cable length, connector to connector (example above is 18/24 meters).
- **X + Y** = used length for connection within system (example above is 4.3 meters).
- **Cable Length - (X + Y)** = available length for conduit run (example above is 13.7 or 19.7 meters).

Illustration 5-6: Cables Group 1 and Ground Cables Group – From Technical Area to Exam Room

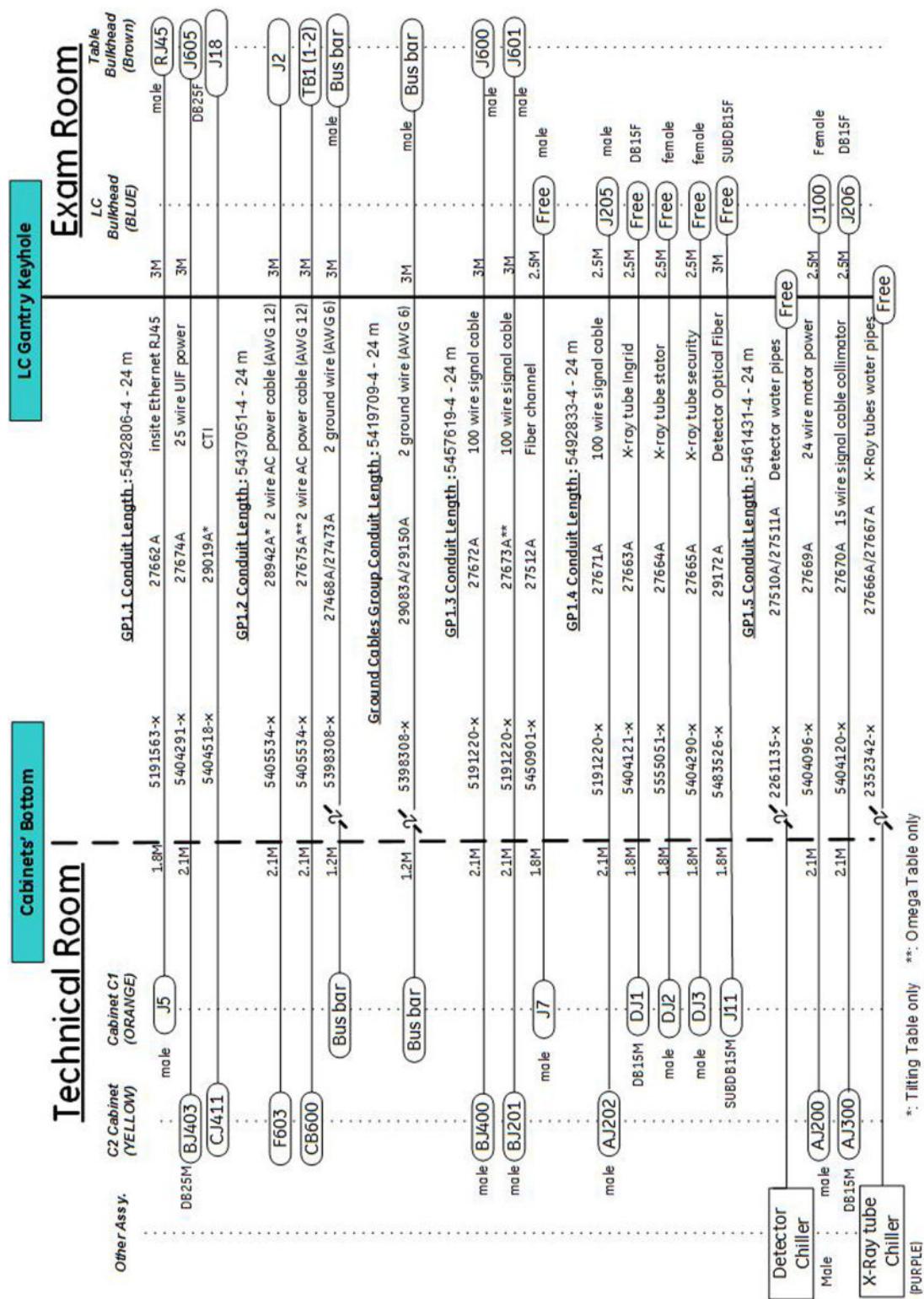


Illustration 5-7: Cables Group 2 – From Technical Area to Control Room

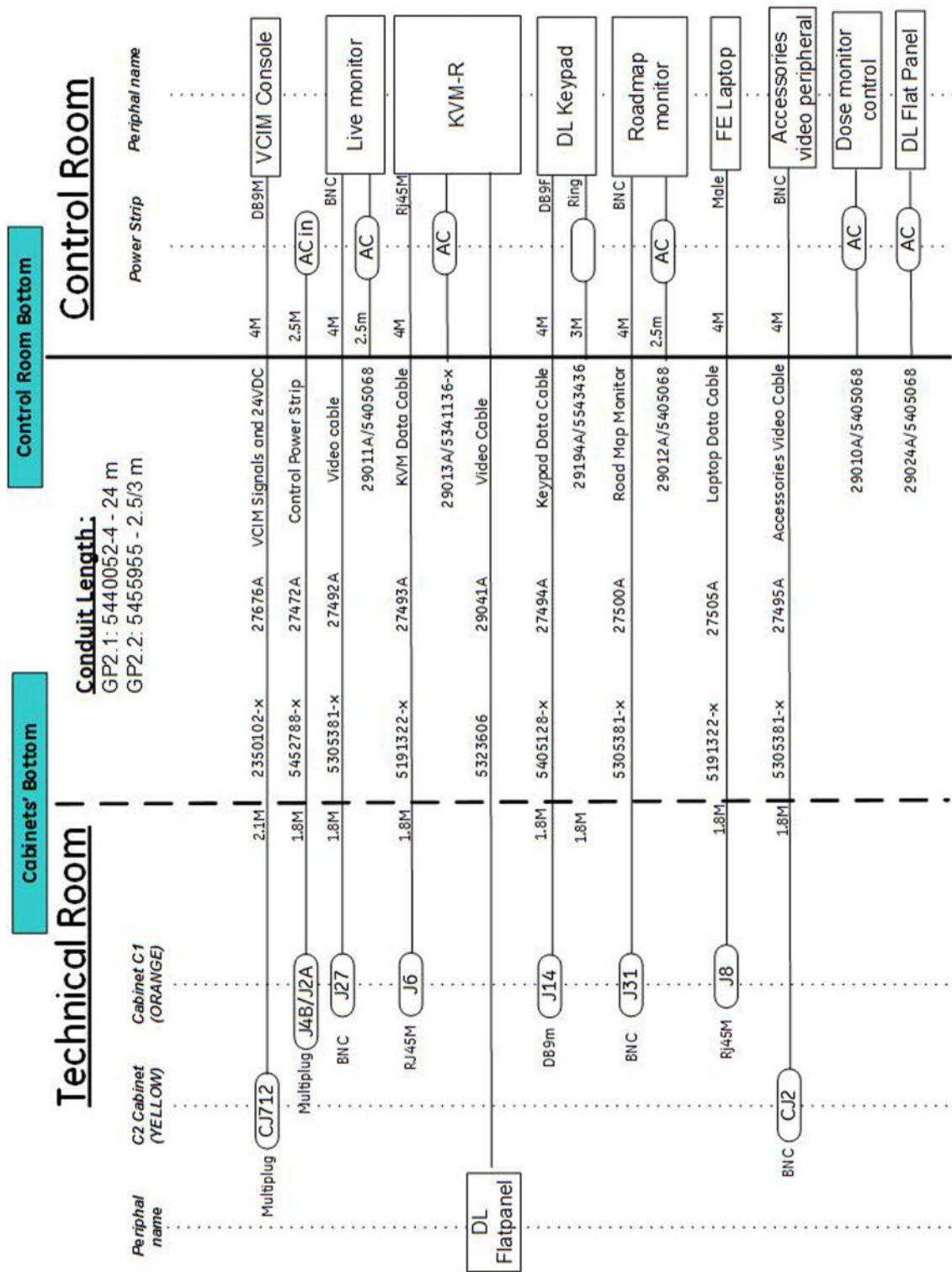


Illustration 5-8: Fast Link Cable Group - Option

Product - Fast Link Cable
5342932 **24 m**

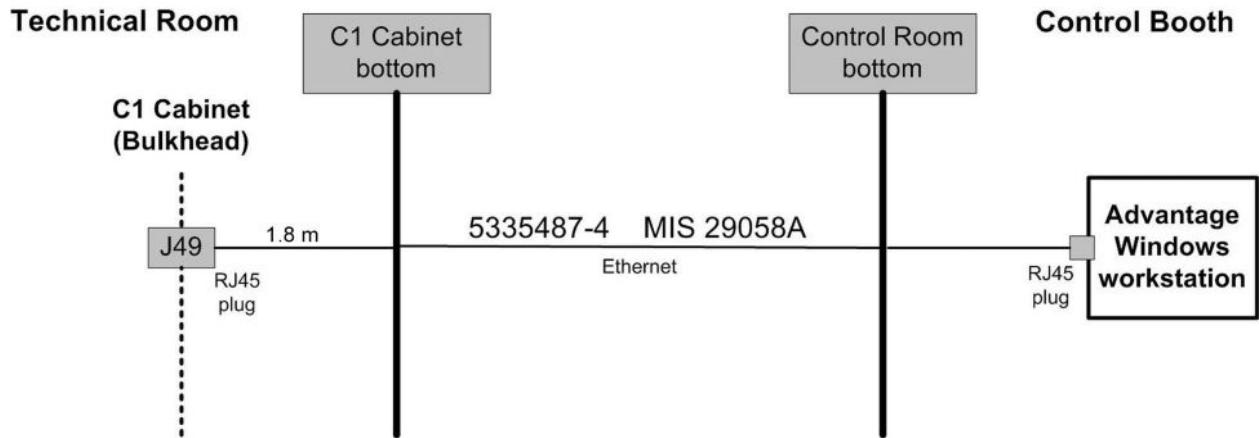


Illustration 5-9: Cables Group 3 – From Technical Area to Technical Area

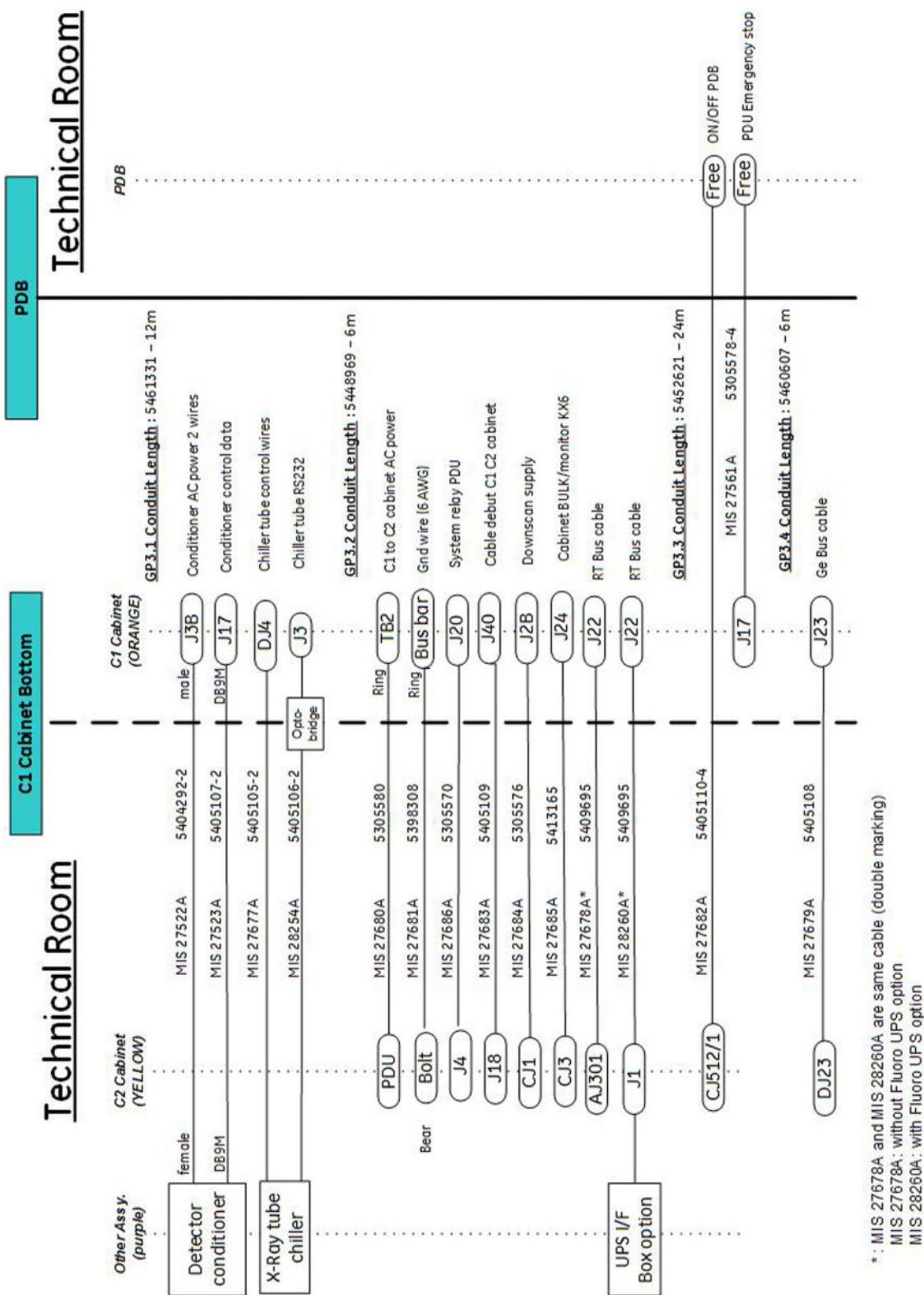


Illustration 5-10: Cable Group 4-2 - From Technical Area to Exam room ceiling

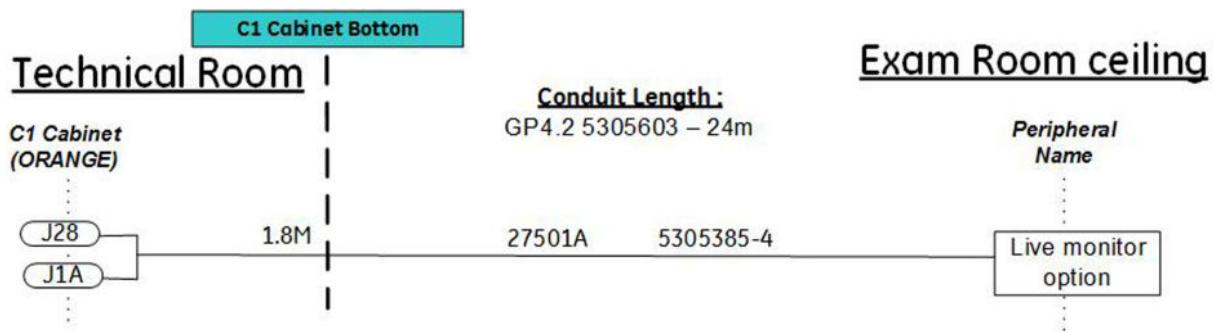
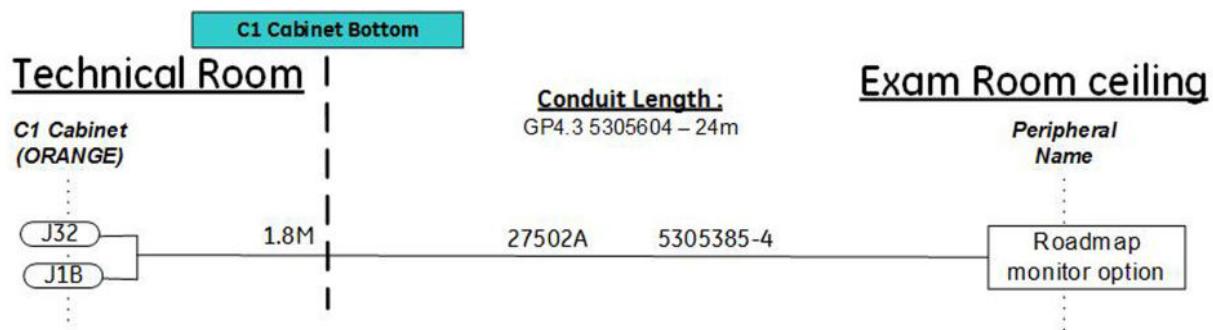


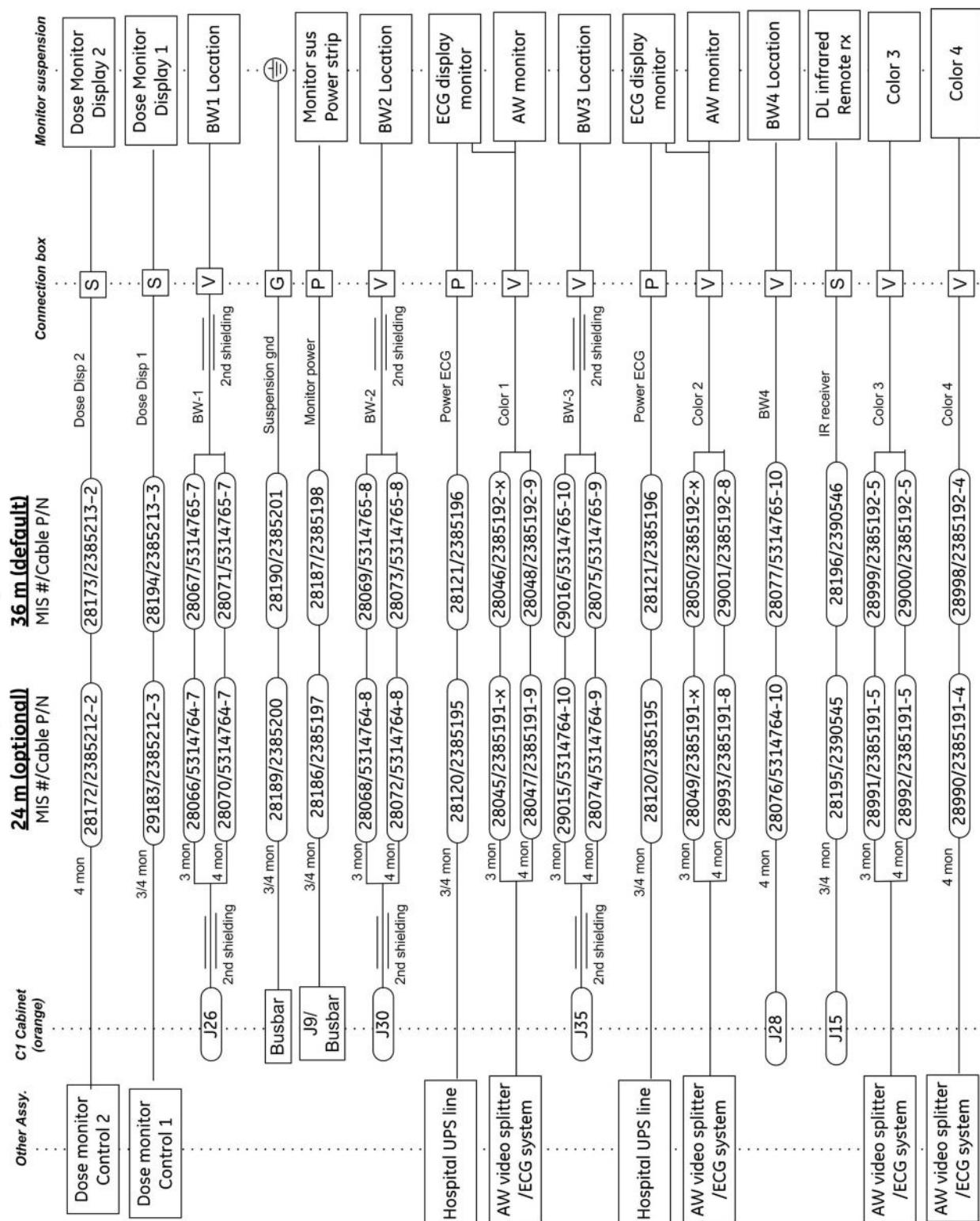
Illustration 5-11: Cable Group 4-3 - From Technical Area to Exam room ceiling



Technical Room

Exam Room Ceiling

OR



For suspension cabling, two length options are available : 24 meter or 36 meter (78 ft 9 in or 118 ft 1 in).

Illustration 5-13: Cable Group 4-5 - From Technical Area to Exam Room ceiling

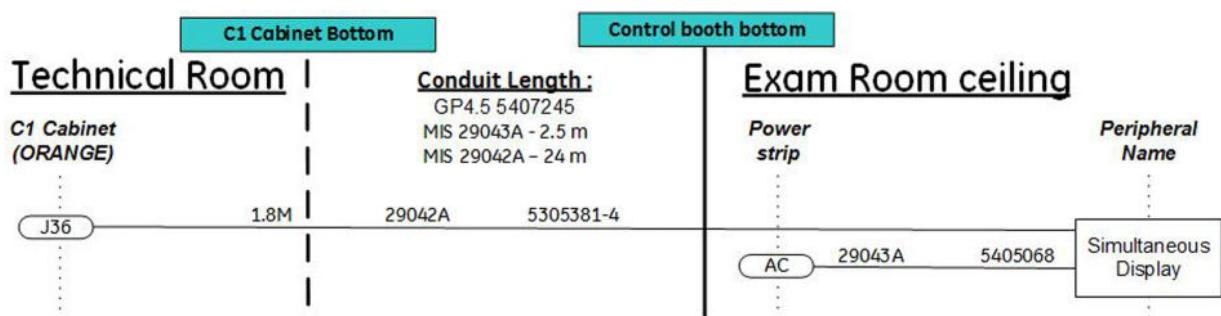
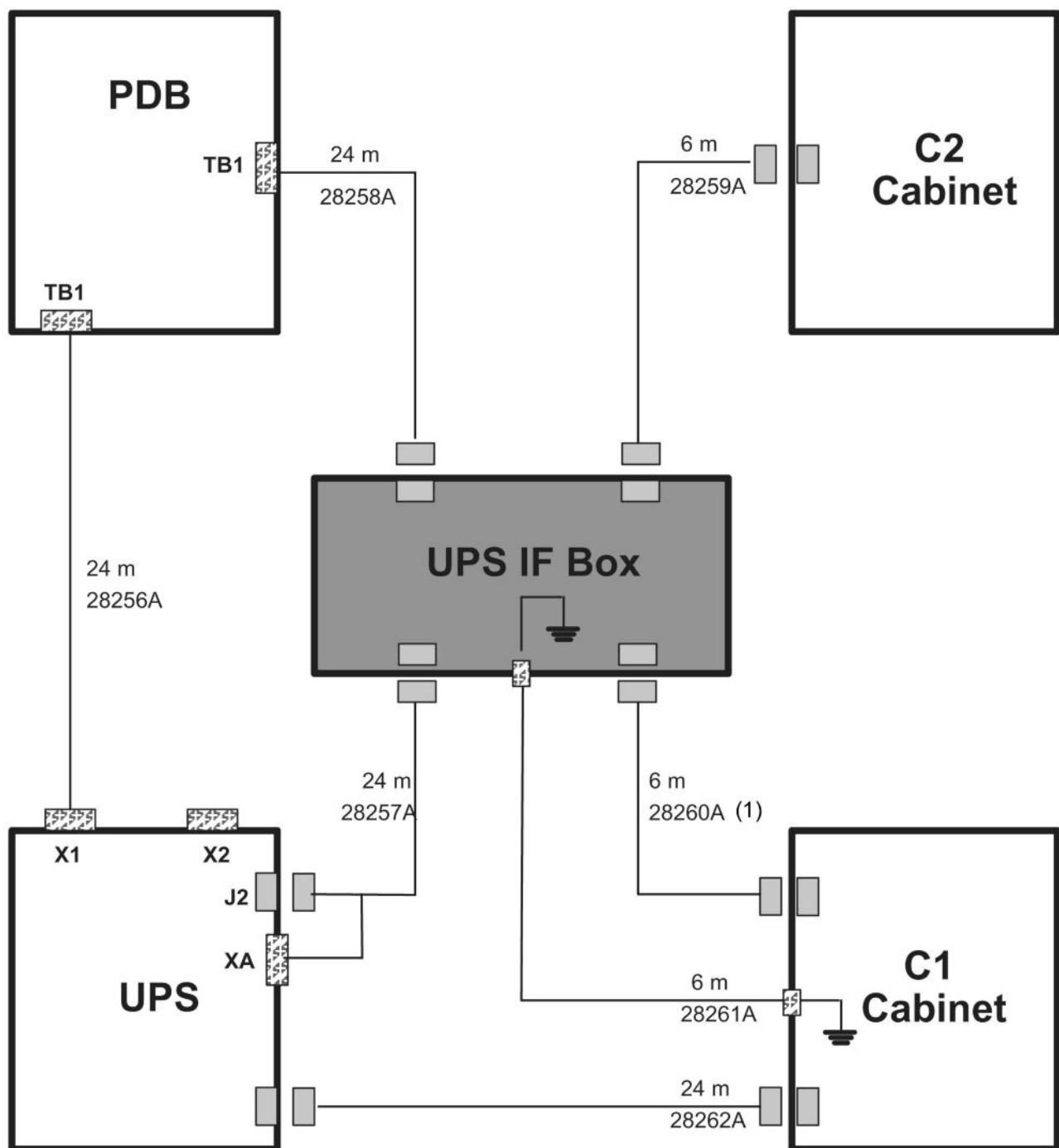


Illustration 5-14: Fluoro UPS Cable Set - Option



NOTE: (1): provided in Group 3 (27678A).

2.1.3 Physical Run - System Core Detail

Table 5-8:

MIS number	Cable Assembly	UL Style	Voltage rating (V)	Max Voltage carried (V)	Cable diameter (mm)	Connector type	Bigger Plug size (mm)
Group n°1 (From C1 / C2 cabinets to Gantry / Table)							
27663A	5404121	2789	600		13.8	DB 11 pin	34.4
27664A	5555051	2463	600		8.3	Metrimate 6 pin	29
27665A	5404290	2463	300		9.2	Metrimate 6 pin	29
27666A	2352342					WATER HOSE	
27667A	2352342					WATER HOSE	
27468A	5398308	1019	600		9.1	Pre-stripping, ring terminal	12
27473A	5398308	1019	600		9.1	Pre-stripping, ring terminal	12
27670A	5404120	2789	300			HES 15 pin	
27510A	2261135					WATER HOSE	
27511A	2261135					WATER HOSE	
27512A	5404122					FIBER OPTIC	
27675A	5405534	Type CL2	150		8.2	Pre-stripping	8.2
27662A	5191563		30		7	AMP 8 pin RJ45	
27669A	5404096	2464	300		16.2	Circular connector	47.8
27671A	5191220	2789	30		10.9	Amplimite 100 pin	84.6
27673A	5191220	2789	30		10.9	Amplimite 100 pin	84.6
27672A	5191220	2789	30		10.9	Amplimite 100 pin	84.6
27674A	5404291	2464	300		10.6	DB 25 pin	56.6
29019A	5404518						
29172A	5483526	2464	300		12.7		
Ground Cables Group							
29083A	5398308	1019	600		9.1	Pre-stripping, ring terminal (only with Tilting table (P/N 5142213-001))	12
29150A	5398308	1019	600		9.1	Pre-stripping, ring terminal (only with Tilting table (P/N 5142213-001))	12
Group no.2 (From C1 / C2 cabinets to Control Booth)							
27676A	2350102		300		8.3	SCSI.50	60
27472A	5452788	4299	300	250		8 plug power strip CEI320 - (HBL4570C plug - ring terminal)	39
27492A	5305381	1354					
27493A	5191322	Category 5	150		5.5	RJ45	14.1
27494A	5405128	2464	300		6	DB 9 pin	30.9

MIS number	Cable Assembly	UL Style	Voltage rating (V)	Max Voltage carried (V)	Cable diameter (mm)	Connector type	Bigger Plug size (mm)
27495A	5305381	1354	30		7	BNC 75 ohms	14.3
27500A	5305381	1354	30		6	BNC 75 ohms	14.4
27505A	5191322	Category 5	150		5.5	RJ45	14.1
29010A	5405068	2464	300			IEC Power Connectors	
29011A	5405068	2464	300			IEC Power Connectors	
29012A	5405068	2464	300			IEC Power Connectors	
29013A	5341136	2464	300			IEC Power Connectors	
29024A	5405068	2464	300			IEC Power Connectors	
29041A	5323606						
29194A	5543436	1015	600		4	IEC Connector - ring terminal	
Optional Fast Link Cable							
29058A	5335487-4				5.9	RJ45	14.1
Group no.3 (From C1 to C2 cabinets)							
27677A	5405105	2463	600		8.3	Metrimate 6 pin	29.8
27678A/ 28260A	5405103	Type CL2	Low Volt-age SCSI		10	SCSI 50C	77
27679A	5405108					Fiber Optic	
27522A	5404292	2464	300		7	CEI 320 - HBL4570C plug	38.6
27523A	5405107						
27680A	5305580			600			
27681A	5398308						
27682A	5405110						
27683A	5405109						
27684A	5305576						
27685A	5413165						
27686A	5305570	2464	300		8.1	DB 9 pin	34.4
28254A	5405106						
27561A	5305578						
Group no.4-2 (From C1 / C2 cabinets to Monitors)							
27501A	5305385	2343	300		16	(CEE22 - ring terminal - BNC50) - (BNC50 - ring terminal - HBL4570C plug)	38.6
Group no.4-3 (From C1 / C2 cabinets to Monitors - Standard Suspension)							
27502A	5305385	2343	300		16	(CEE22 - ring terminal - BNC50) - (BNC50 - ring terminal - HBL4570C plug)	38.6
Group no. 4-4 (LCD Suspension)							
28045	2385191		30V			VGA	
28046	2385192		30V			VGA	

MIS number	Cable Assembly	UL Style	Voltage rating (V)	Max Voltage carried (V)	Cable diameter (mm)	Connector type	Bigger Plug size (mm)
28047	2385191		30V			VGA	
28048	2385192		30V			VGA	
28049	2385191		30V			VGA	
28050	2385192		30V			VGA	
28051	2385191						
28052	2385192						
28053	2385191						
28054	2385192						
28066	5314764	1354	30V			BNC	14.4
28067	5314765	1354	30V			BNC	14.4
28068	5314764	1354	30V			BNC	14.4
28069	5314765	1354	30V			BNC	14.4
28070	5314764	1354	30V			BNC	14.4
28071	5314765	1354	30V			BNC	14.4
28072	5314764	1354	30V			BNC	14.4
28073	5314765	1354	30V			BNC	14.4
28074	5314764	1354	30V			BNC	14.4
28075	5314765	1354	30V			BNC	14.4
28076	5314764	1354	30V			BNC	14.4
28077	5314765	1354	30V			BNC	14.4
28078	5314764						
28079	5314765						
28080	5314764						
28081	5314765						
28082	5314764						
28083	5314765						
28084	5314764						
28085	5314765						
28086	5314764						
28087	5314765						
28088	5314764						
28089	5314765						
28120	2385195		300			Power	
28121	2385196		300			Power	
28172	2385212	2789	30			DB-9pin	
28173	2385213	2789	30			DB-9pin	
28186	2385197		300			Power	

MIS number	Cable Assembly	UL Style	Voltage rating (V)	Max Voltage carried (V)	Cable diameter (mm)	Connector type	Bigger Plug size (mm)
28187	2385198		300			Power	
28189	2385200					G/Y	
28190	2385201					G/Y	
28194	2385213	2560	30		6.6		
28195	2390545	2789	300			DB-9pin	
28196	2390546	2789	300			DB-9pin	
28990	2385191						
28991	2385191						
28992	2385191						
28993	2385191						
28998	2385192						
28999	2385192						
29000	2385192						
29001	2385192						
29015	5314764						
29016	5314765						
29183	2385212	2560	30		6.6		
Group no. 4-5							
29042A	5305381						
29043A	5405068						
Fluoro UPS Option Cable Set							
28256A	5139650						
28258A	5139792						
28262A	5191563						
28257A	5305596						
28261A	5398308						
28259A	5405103						

2.2 MIS (Master Interconnect System)

Innova system interconnect cables are described in MIS (Master Interconnect System) documents. These documents specify all interconnections between components within the system.

Reference: For specific Vascular system interconnect maps and connection details, refer to the following

- *System MIS Map*
- *System MIS Charts*

General Guidelines

Innova System introduce a new system interconnect with a star distribution for all cables from the technical area. The cables are shipped on spools to create cable groups. Cable group 1 for Exam room and cable group 2 for control room. The cable group shall be put in place during the same action. The cables are routed in the same duct.

The HV cables could be pulled separately.

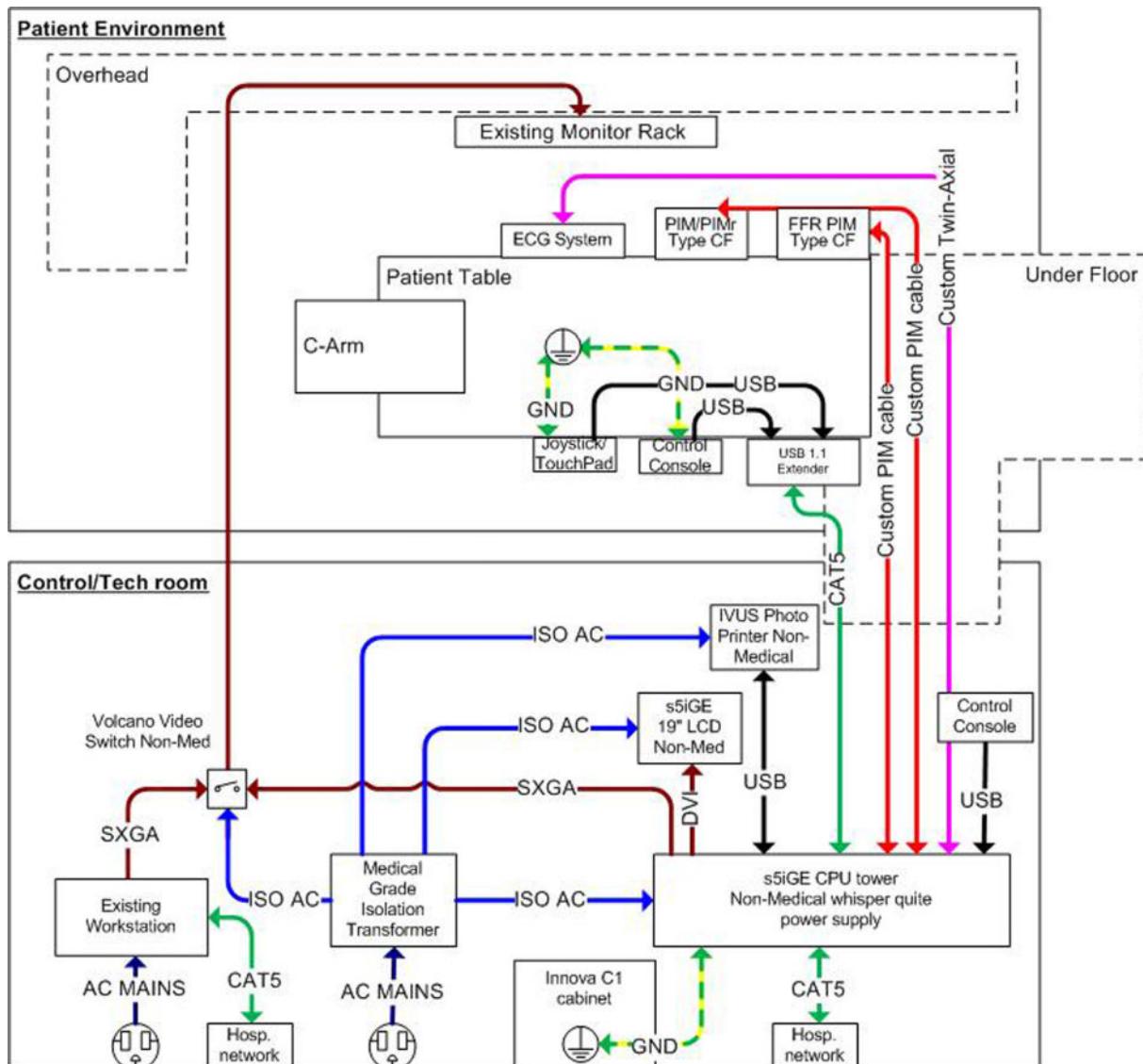
2.3 IVUS Wiring

2.3.1 For IVUS rev 2 wiring

The connection cables between the IVUS Control Room and Procedures Room components run in a dedicated under floor conduit (see [Cable Channelling](#)).

A GND wire between the C1 cabinet and the IVUS Rev 2 CPU runs in the underfloor conduit prepared for the Control Booth cables between Equipment Room and Control Room.

Illustration 5-15: IVUS rev 2 wiring

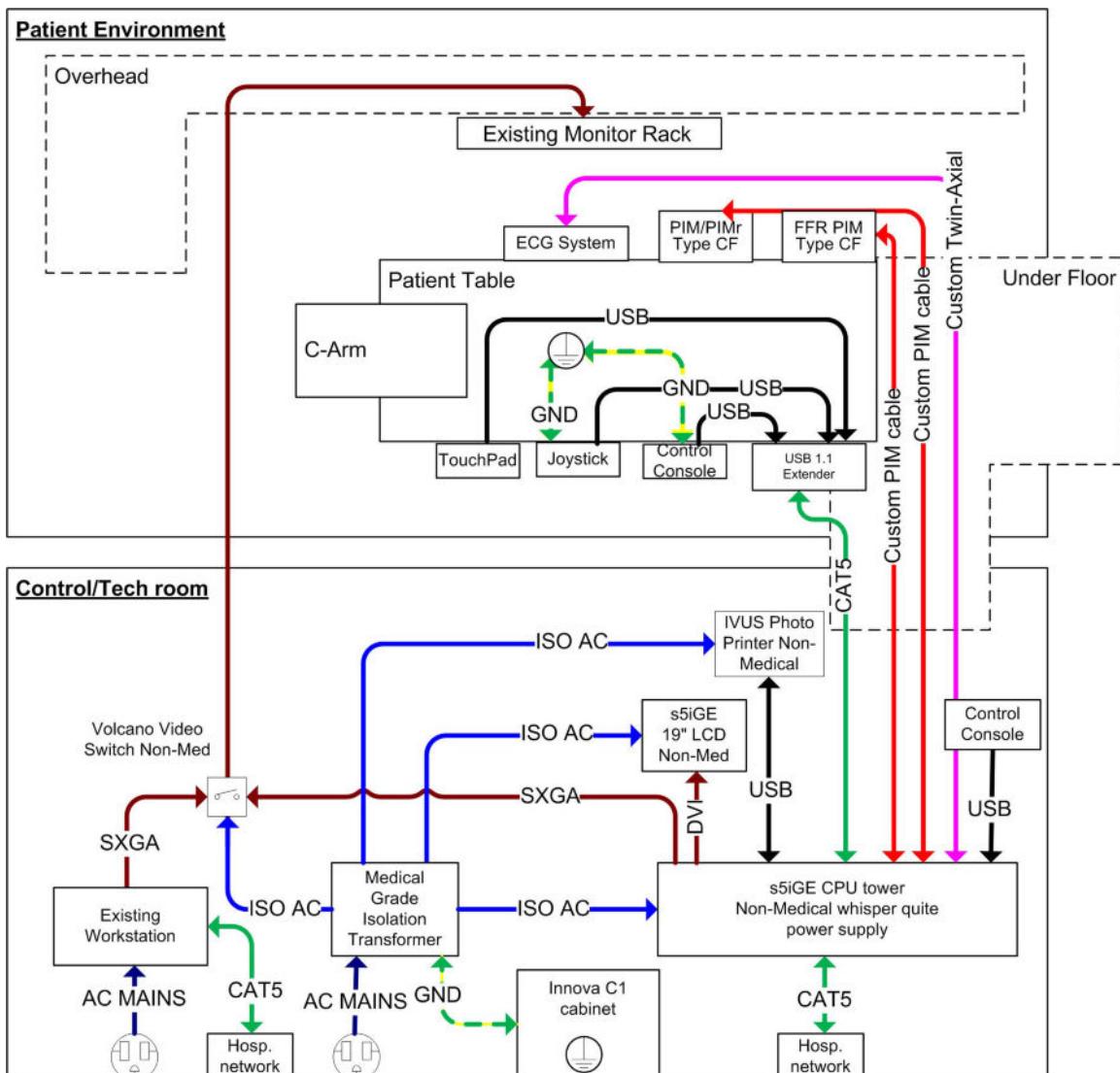


2.3.2 For S5I GE rev 2+ wiring

The connection cables between the S5I GE Control Room and Procedures Room components run in a dedicated under floor conduit (see [Cable Channeling](#)).

A GND wire between the C1 cabinet and the S5I GE Rev 2+ Isolation Transformer runs in the underfloor conduit prepared for the Control Booth cables between Equipment Room and Control Room..

Illustration 5-16: S5I GE rev 2+ wiring



2.3.3 For IVUS rev 3 wiring

Refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

2.4 Cable Channeling

2.4.1 General

High voltage and power cables must be separated from other cables. Use a separate trough in the duct system, or use a separate conduit. Minimize cable length between the line disconnect and the System Cabinet power unit to reduce voltage regulation problems and wiring costs.

For information about the cables supplied with your system, please refer to [Physical Runs](#).

2.4.2 Conduit

Separate conduits must be used for power and signal wires. These wires must be kept separated from each other.

Using conduit imposes some important considerations when used with this system. Of primary concern, the majority of cables used are pre-terminated. Pre-termination greatly simplifies interconnection but makes cable-pulling difficult because of the added dimensions of the connectors.

Conduit must be large enough to pass the cable and connector through with all other cables already in the conduit. Also, the size of conduit chosen must allow for future growth. There is the possibility of additional cables being added later as the system is developed and options are added.

The use of conduit is recommended for cables running overhead between rooms, especially when a diagonal run provides the shortest cable path

Separate conduits must be used for Hospital and Fluoro UPS power wires. These wires must be kept separated from each other (at least 30 cm).

2.4.3 Electrical Ducts

NOTE: (For IVUS Rev 3 option) refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

It's important that electrical ducts have separate compartments for power and signal wires. These wires must be kept separated from each other for proper system operation.

Electrical ducts have advantages, when used with a single room or two adjacent rooms. Electrical ducts combine cabling in a neat and functional appearance, with accessibility and room for expansion.

NOTE: Medrad AVANTA, IVUS and Mac-lab cables exit behind the table in the patient room.

NOTE: For **Fast Link** cable (C1 cabinet - AW station), the static operation bending radius must be at least 4 times the outer cable diameter.

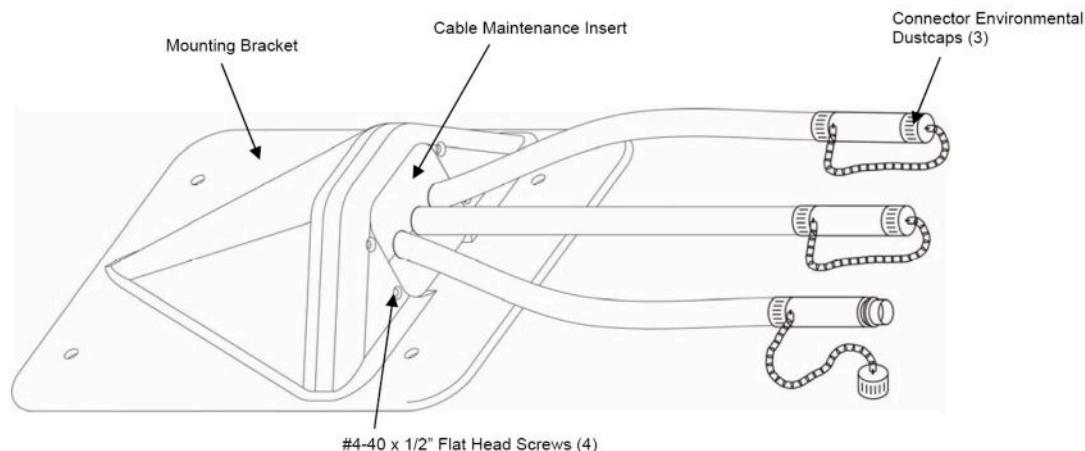
It is the responsibility of the site planner to provide the appropriate solution to the table exit (e.g gas box, Clab II, Tram module, connection interface box)

NOTE: Specific Recommendations for installation with GE ECG Device such as MacLab, CardioLab or ComboLab:

- TRAM RAC in Exam Room with cable 2016134-106 routed back to Control Room where the other modules & PC are installed
- If no GE Maclab cable 2016134-106 installed between the TRAM (Exam Room) and the Control Room, need to route it so that installation/connection of Physio module can be made in Control Room.

NOTE: MEDRAD Avanta Table mount with floor Medrad bracket (this configuration is not authorized with an Tilting table): A 76.2 mm (3 in) and max 25 m (984 in) length conduit between technical room and patient room shall be prepared below the floor for the three injector cables. It is recommended to use the MEDRAD Avanta floor mounting bracket to cover the duct hole in the patient room if there is no gases box.

Illustration 5-17: MEDRAD Avanta mounting bracket



Floor mount installation can be accomplished one of two ways:

- Connectors mounted in trough under mounting bracket (Figure 1)
- Connectors mounted above mounting bracket (Figure 2)

Illustration 5-18: MEDRAD Avanta floor mounting methods

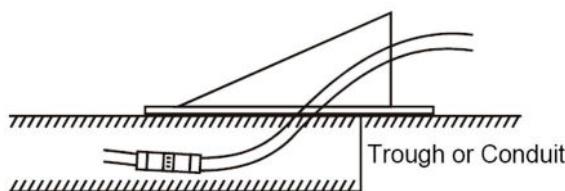


Figure 1

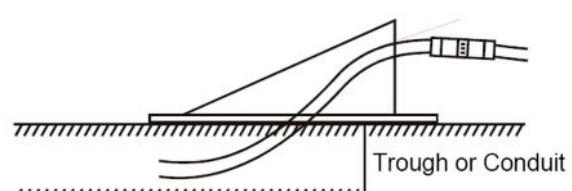
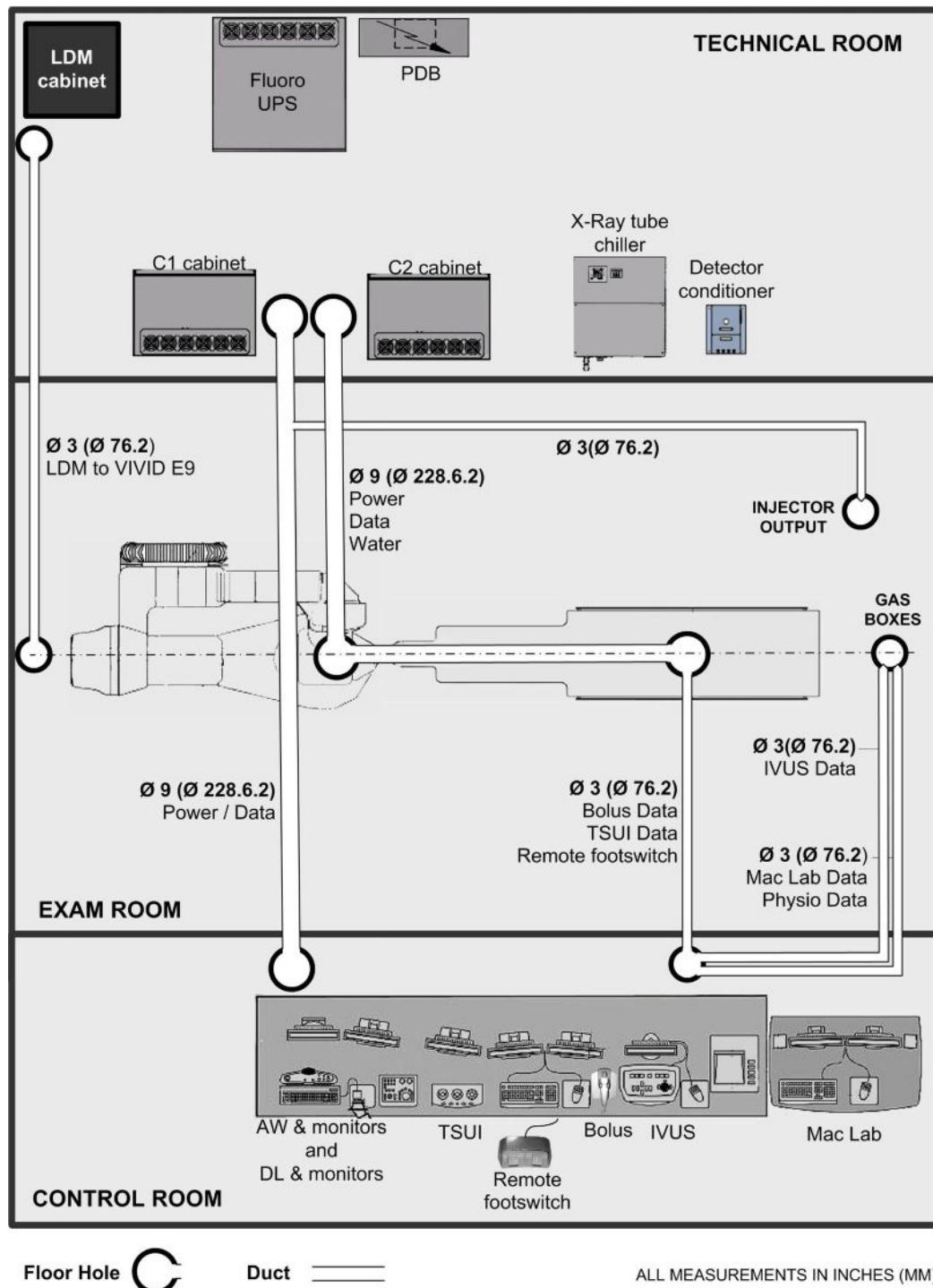


Figure 2

For further MEDRAD Avanta floor mounting, see the Installation guide MEDRAD Avanta Floor Mounting Bracket.

Illustration 5-19: Cable Channeling Layout



NOTE: 18 meters (59 ft) is the only cable length available for the Remote TSUI box data cable connecting remote TSUIs in the control booth and the patient table.

NOTE: Never connect the **Physio** module to the ground.

The Physio cable can run in the same conduit as the Bolus cable. In this case, it is required to have a conduit between the table and the physio gases box.

If no conduit available between rear of table and Control Room (no Remote TSUI, no IVUS, no MacLab...), need to define proper cable routing or create new conduit as per PIM requirements.

If there is no physio gases box behind the table in the lay out, find a local solution to hide the hole in the floor and the cable exit.

On some sites there is only one conduit between Exam room and Control room for the IVUS & MacLab.

2.4.4 IVUS S5I Option

2.4.4.1 Floor Conduits

NOTE: (For IVUS Rev 3 option) refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

A 3 inches and max. 25 m long conduit between Control and Patient Rooms shall be prepared below the floor for the S5I GE Innova interconnection cables.

- Locate the position of the S5I GE components in the Control Room, according to the workflow requirement of the operators.
- Establish a 76 mm (3") conduit below floor from the S5I GE CPU location in the Control Room, to the foot end area of the Patient Table in the Procedures Room:
 - Locate the Procedures Room exit of the conduit close to the centerline of the Patient Table
 - Locate it as close to the Patient Table base as possible, considering :
 - Room for access to the table for service purposes
 - Limitation from other cables connected to the table
 - Location of the table plate
 - Location of Mac-Lab conduit
 - Location of Physio or Med Gases connections
 - The S5I GE cables are moving cables, they have to allow the Patient Table motions (Panning, Lifting, Tilting and Rotating)
- Use similar solutions (i.e. protection against cleaning fluids, etc.) as it is used for the Mac-Lab conduit
- See also guidance on Typical Layout drawing 4-58f sheets E1 and S2, showing Mac-Lab and P/M Gas location recommendations.

Estimated labour time: 1 hour

2.4.4.2 Run Interconnection cables through the Conduit

2.4.4.2.1 For IVUS Rev 2 option

Use the IVUS Rev 2 PIM (#806365002) as reference to the procedures below.

- Unpack , and check the completeness of the Preinstallation kit :
 - 1 pcs - Custom PIM cable 30 m (98 ft, 5 in) - Volcano ID :806452001
 - 1 pcs - CAT5 cable 30 m (98 ft, 5 in) - Volcano ID :807055001
 - 1 pcs - Custom Twin axial cable - Volcano ID :806380001
 - 1 pcs - Ground Cable, 30 m (98 ft, 5 in) s5i – Volcano ID :806889001
 - 1 pcs - IVUS Rev 2 Preinstallation Manual - Volcano ID :806365002
- Install 806889001 Ground Cable through the Control Booth conduit (see note)
- Install the cables through the IVUS conduit from the Control Room end :
 - 806452001 PIM cable, round shaped connector end to the Procedures Room
 - 807055001 - Ethernet cable (1 pcs)
- Install the optional cables through the IVUS conduit from the Control Room end :
 - 806380001 - ECG cable in case, if NON Mac-Lab ECG will be installed
 - 808631002 – FFR cable option - 15m
 - 808632002 – FFR cable option - 27m

NOTE: For the ground cable, see the IVUS Rev 2 service manual for special instructions concerning cable running and cutting .

NOTE: If the installation has the **Mac-Lab ECG** in the configuration, the ECG cable shall not be run through the conduit, it will be used in the Control Room **to connect Mac-Lab and IVUS PCs**.

NOTE: You should consider running the ECG cable through the conduit in case, the customer requires interconnecting to a **NON Mac-Lab ECG** at the table base.

In this case you **shall hide the unconnected cable in the conduit** for future expansion options.

NOTE: Same applies to the 2nd Ethernet cable, if a later extension of the IVUS remote controls expected.

- **Ensure sufficient cable length on the Procedures Room end of the cables according to the table below**

Cable	Free length from the conduit exit
IVUS PIM cable *	2.4 m (8 ')
FFR cables for Pimetted **	0m (0')

Cable	Free length from the conduit exit
Control Station cable	2.1 m (7 ')
Joystick cable	2.7 m (9 ')
ECG cable	Depends on the ECG location

* Consider the 12' length of the PIMr cable, if PIMr exists.

** Pimette has 3 m cable.

NOTE: The incorrect cable length might cause reliability issues during the operation !

Make sure, that the out of conduit parts of the cables with the connectors are properly protected during the rest of the installation

2.4.4.2.2 For S5I GE Rev 2+ option

Use the S5I GE Rev 2+ PIM (#806365026) as reference to the procedures below.

- Unpack , and check the completeness of the Preinstallation kit:
 - 1 pcs - Custom PIM cable 27 m (88.2 ft) - Volcano ID: 806452008
 - 1 pcs - CAT5 cable 30 m (98 ft, 5 in) - Volcano ID: 807055001
 - 1 pcs - Custom Twin axial cable - Volcano ID: 806380001
 - 1 pcs - Ground Cable, 30 m (98 ft, 5 in) s5i – Volcano ID: 806889001
 - 1 pcs - S5I GE Rev 2+ Preinstallation Manual - Volcano ID: 806365007
 - 1pcs – FFR cable 27 m - Volcano ID: 808632001
- Install 806889001 Ground Cable through the Control Booth conduit (see note)
- Install the cables through the S5I GE conduit from the Control Room end :
 - 806452008 PIM cable, round shaped connector end to the Procedures Room
 - 807055001 - Ethernet cable (1 pcs)
 - 808632001 – FFR cable - 27 m
- Install the optional cables through the S5I GE conduit from the Control Room end: 806380001 - ECG cable in case, if NON Mac-Lab ECG will be installed

NOTE: For the ground cable, see the S5I GE Rev 2+ service manual for special instructions concerning cable running and cutting .

NOTE: If the installation has the **Mac-Lab ECG** in the configuration, the ECG cable shall not be run through the conduit, it will be used in the Control Room **to connect Mac-Lab and S5I GE PCs**.

NOTE: You should consider running the ECG cable through the conduit in case, the customer requires interconnecting to a **NON Mac-Lab ECG** at the table base.

In this case you **shall hide the unconnected cable in the conduit** for future expansion options.

- Ensure sufficient cable length on the Procedures Room end of the cables according to the table below

Cable	Free length from the conduit exit
S5I GE PIM cable *	2.4 m (8')
FFR cable for Pimette **	0 m (0')
Control Station cable	2.1 m (7')
Joystick cable	2.7 m (9')
ECG cable	Depends on the ECG location

* Consider the 3 m length of the short cable, if the split cable preinstallation exists.

** Pimette has 3 m cable.

NOTE: The incorrect cable length might cause reliability issues during the operation !

Make sure that the out of conduit parts of the cables with the connectors are properly protected during the rest of the installation.

2.4.4.2.3 For IVUS Rev 3 option

Refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

2.4.5 Vivid E9 Ultrasound Option

Vivid E9 connection with LDM needs a conduit between the Technical Room (LDM cabinet) and the Vivid E9 cables access point on the head end wall, close to the room floor and aligned with the Patient table centerline (see [Illustration 5-19](#)).

In case if there is an existing floor conduit for the LDM linkset cables with an exit close to the Patient Room wall, it is possible to use this duct for the Vivid E9 LDM image link, and continue with a wall conduit to reach to the Vivid E9 cables access point on the head end wall of the Patient Room.

For Vivid E9 power connection preparation, please refer to *VIVID E9 Service Manual GA091568* Electrical requirements section.

For Vivid E9 network connection preparation, please refer to *VIVID E9 Service Manual GA091568* Insite/Network Connection section.

3 Lighting Specifications

3.1 Room Light Distribution

3.1.1 Requirements for lighting

Requirement for lighting concern the following, general, light-technique characteristics:

- Illuminator level.
- Lighting distribution.
- Preventing the operator from being dazzled by the light (by direct light sources or by reflection on bright objects).

The illumination level must be compliant with established lighting technical rules and be as constant as possible.

Technical room, operating room and control room shall be provided with appropriate lighting in the maintenance area (maintenance area to be considered are service workplaces). It corresponds to service areas as defined for any of the product components.

The minimum required average luminance E_m shall be of 500Lx and minimum color rendering factor R_a of 80 as per IEC/EN 12464-1 (Light and lighting. Lighting of work places. Indoor work places: Illumination requirements for indoor workplaces corresponding to assembly of medium size electrical components, e.g. control panel) for the electrical industry).

3.1.2 Lighting Relay

Innova System has the ability to control an external relay that applies power to the room light (dry contacts).

The relay is to be provided by the hospital or contractor.

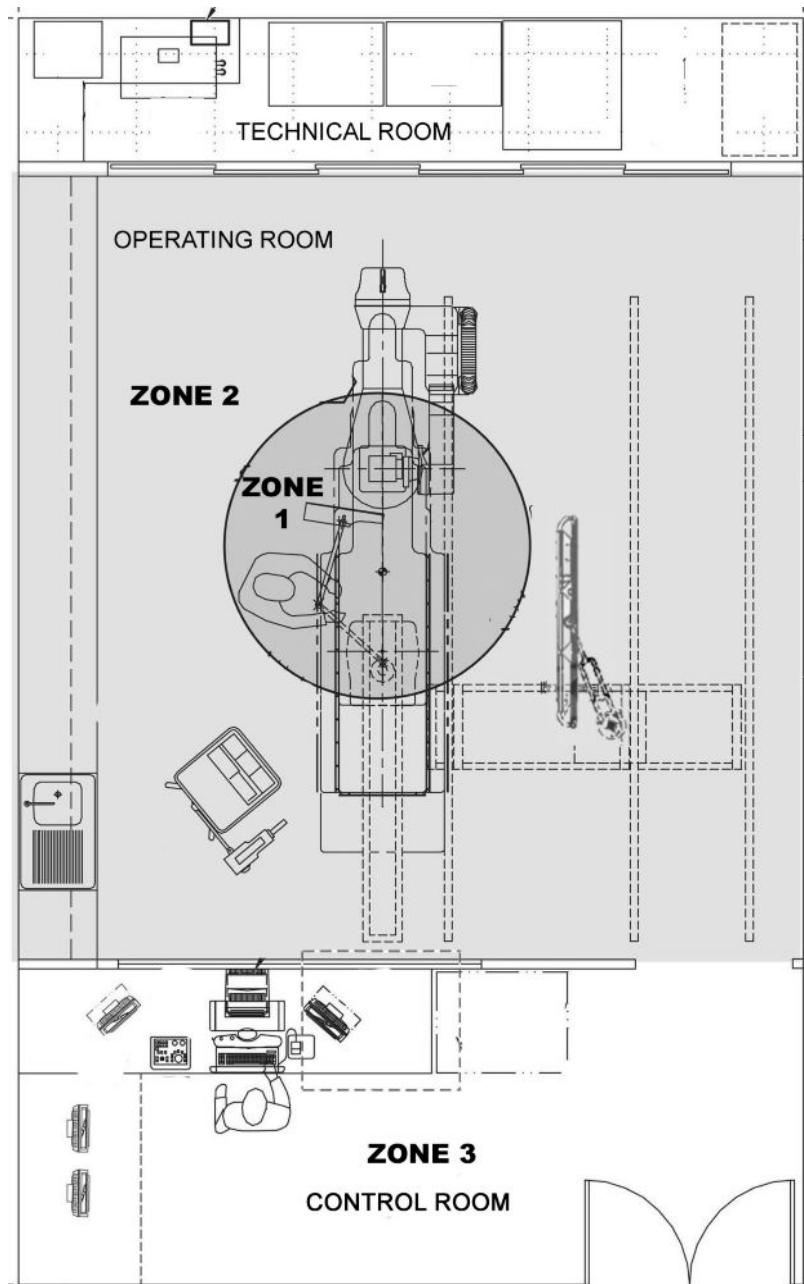
The wire size to connect to the C2 cabinet is 1.5 mm² (AWG 14).

Relay rating is max 600 VAC

The room light distribution is wired between the C2 cabinet and the PDB.

3.1.3 Room Lighting for System

Illustration 5-20: Room Lighting Zones



3.1.4 Windows and curtains

When the examination room has a window with an aperture outside of the controlled light area (day light, other...) a curtain has to maintain the light intensity under a limit fixed to 150 lux.

Chapter 6 Communication Requirements

1 Network Requirements

1.1 Insite/Network Connection

The preferred Insite connection uses a broadband modem. This connection requires a dedicated Ethernet Jack (RJ45) that must be located less than 1 meter (3 feet) from the C1 cabinet.

For complete descriptions of these connectivity solutions, please refer to the Broadband Solutions catalogue available through your local GEHC sales and service representative.

Connectivity Process and pre-installations checklists are available in the Broadband Connectivity PIM available through your local GEHC sales and service representative.

InSite requires an Internet Address connecting it to the Innova System. This address must be available before installing the system. A request form has been defined. For more information, please refer to [IP Addressing Process](#) or contact your GEMS OLC representative.



NOTICE

The C1 cabinet comes equipped with a Firewall unit. The hospital network must be capable of connecting to this firewall. In the case that it cannot be, please contact GE Healthcare to discuss alternatives.

1.2 Options

1.2.1 IVUS Ethernet Network Requirement

NOTE: For IVUS Rev 3 option, refer to [Volcano s5i Imaging System Integration in IGS Systems - Service Manual](#).

The S5I GE CPU located in the control room shall be connected to the hospital Ethernet network. A wall Ethernet outlet shall be available in the control room to connect the device.

DICOM Image Storage:

Saving patient cases to DVD: The archived images are stored in DICOM format with Volcano s5i system acting as a File Set Creator (FSC), following the guidelines in the 2004 DICOM 3.0 specification.

Sending patient cases to DICOM server: The Volcano s5i supports the ultrasound multi-frame image storage SOP class as an SCU (service class user).

For more information, see the Volcano s5 DICOM Conformance Statement located in the Volcano Service Manual.

1.2.2 Vivid E9 Ultrasound Communication Requirements

1.2.2.1 VIVID E9 Supported networks

10/100/1000 Mbit Ethernet/DICOM network (option)

1.2.2.2 VIVID E9 InSite Requirements

InSite requires an Ethernet connection either via:

- 10/100 Mbit

or

- 10/100/1000 Mbit Interface

1.2.2.3 VIVID E9 Purpose of the DICOM network function:

DICOM services provide the operator with clinically useful features for moving images and patient information over a hospital network. Examples of DICOM services include the transfer of images to workstations for viewing or transferring images to remote printers. As an added benefit, transferring images in this manner frees up the on-board monitor and peripherals, enabling viewing to be done while scanning continues. With DICOM, images can be archived, stored, and retrieved faster, easier, and at a lower cost.

For further details please refer *VIVID E9 Service Manual GA091568TPH* Section 2-3-6.

1.2.2.4 VIVID E9 IP Addressing Process

Please refer *VIVID E9 Service Manual GA091568TPH* Section 2-3-6.

2 IP Addressing Process

To obtain an IP address, contact the following for your pole:

- **GEMSAM:**

Contact: OnLine Center–Americas, Network Products and Services (NP&S)

Telephone: 1–800–321–7937

NOTE: Press [1] for the Online Center. Follow the phone tree instructions to select X-Ray modality. When prompted, select the option for obtaining an IP address.

- **GEMSE:**

Use the new mail form called */NSFORM.xls* or */NSFORM.txt* for obtaining an IP Address.

If you have questions or need clarification regarding the use of this form, do not hesitate to ask the Operation support OnLine.

Contact: OnLine Center–Europe

Telephone: +33 (0)1 30 83 13 00

FAX: +33 (0)1 30 70 99 70

NOTE: The INSITE FORM is on the formatted sheet (.xls) or text sheet (.txt) that can be found on the Service CD–Rom.

- **GEMSA:**

Contact: OnLine Center–Asia

Network Products and Services (NP&S)

Telephone: (81) 426 56 0033

FAX: (81) 426 56 0053

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Chapter 7 Appendix

1 PIST0021 - US Fluoro UPS and PDB connection

1.1 Personnel Requirements

Personnel Requirements	Preliminary Reqs	Procedure	Finalization
1	Not Applicable	1 hour	Not Applicable

1.2 Preliminary Requirements

1.2.1 Tools and Test Equipment

Item	Qty	Effectivity	Part#	Manufacturer
Standard Service Engineer's toolkit	1	-	-	-

1.2.2 Required Conditions

Condition	Reference	Effectivity
The PDB is already installed	-	-
Power cables are as follows: (1) From PDB to UPS rectifier input = 4x6 AWG recommended (4x3 AWG max) (3ph + PE). (2) Bypass = 5x8 AWG recommended (5x3 AWG max)(3ph + N +PE). (3) From UPS output to PDB = 5x8 AWG recommended (5x3 AWG max)(3ph + N+PE)..	-	-
Cable sizing shall follow local requirements.	-	-
Power cable Mechanical Terminals Torque Specification: < 3 Nm / < 27 lb.in for 20 AWG to 3 AWG.	-	-
Check the UPS shipping date marked on the packaging label. Commissioning of the UPS must be performed within 3 months after that date.	-	-

1.3 Procedure

1.3.1 Connection of the Power Distribution Box

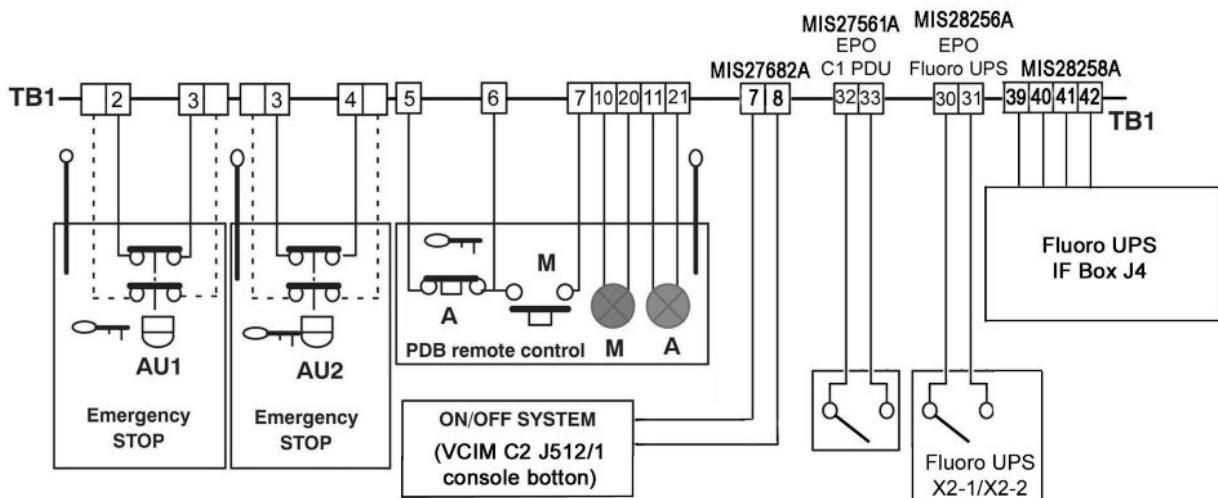


NOTICE

Hospital main power must enter from the Top. Correct phase sequence during installation is very important. Phase failure relays are provided to monitor voltages less than 85%, loss of a phase, or incorrect phase sequence. Corrections to the phase sequence must be corrected only at the line side of the main circuit breaker CB1.

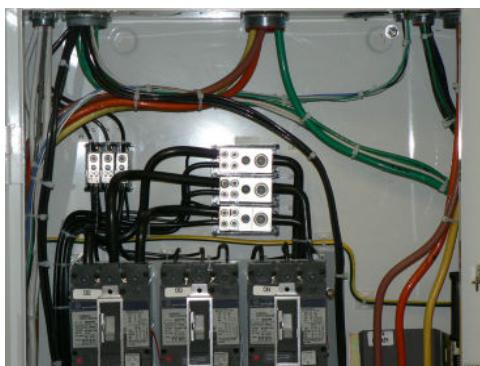
1. Connect the ON/OFF, Remote control and Emergency Power Off devices.

Illustration 7-1:



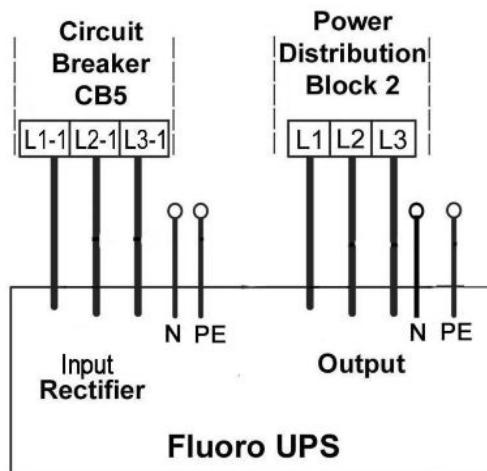
2. If the Fluoro UPS option is installed, remove the L1A, L2A and L3A jumpers linking the Power Distribution Block 1 and the Power Distribution Block 2.

Illustration 7-2:



3. Connect the Power distribution cables.

Illustration 7-3:

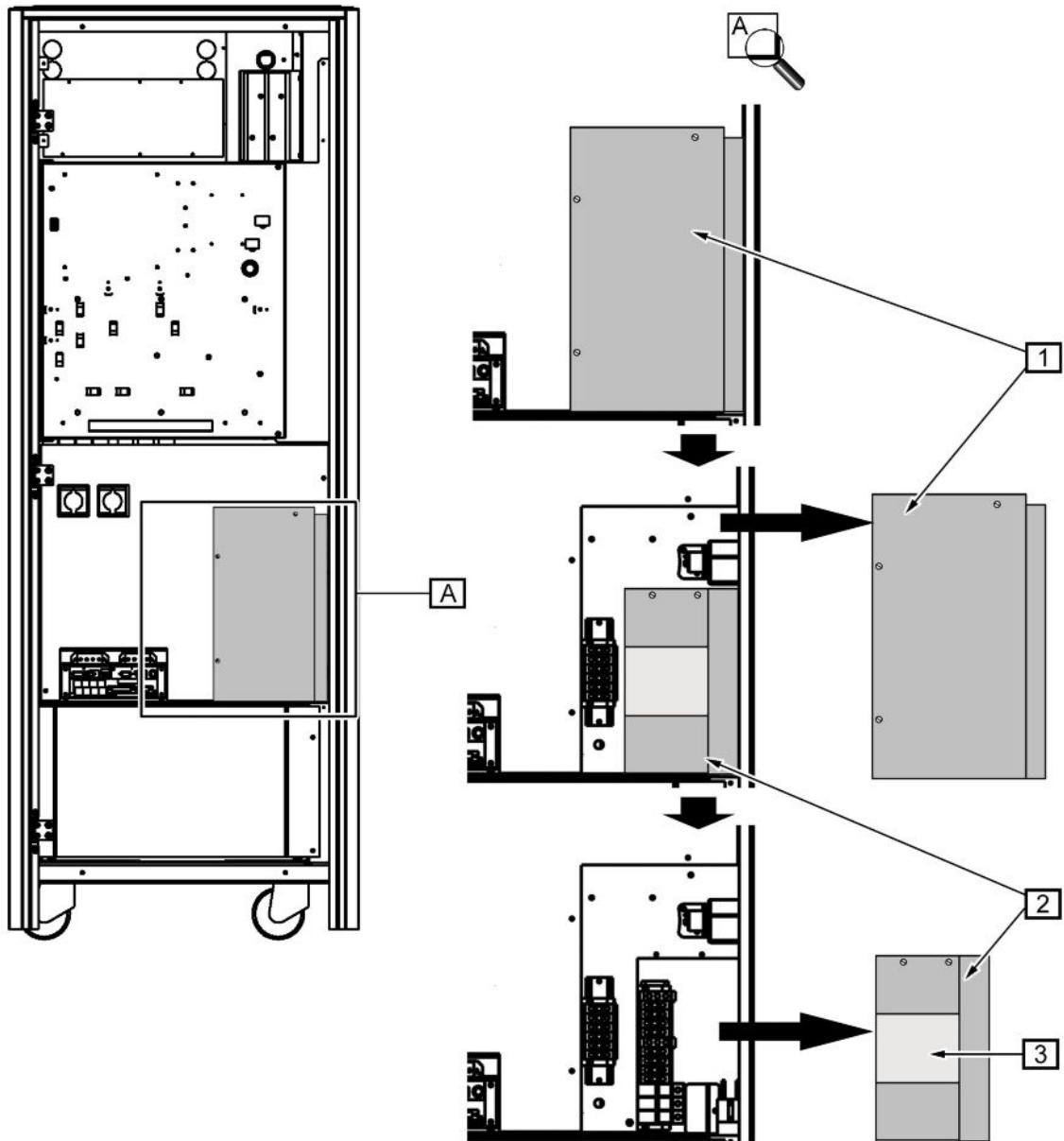


1.3.2 Connection of the Fluoro UPS Option Cables

1. Connect the Fluoro UPS power cables.

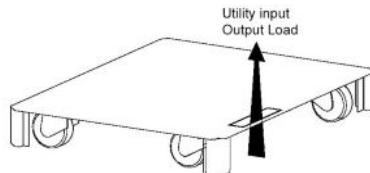
- a. Remove the protection panel **1**.

Illustration 7-4:



- b. Remove the protection panel (**2**, [Illustration 7-4](#)).
- c. Remove the metallic window **3** from the protection panel **2** ([Illustration 7-4](#)).
- d. The power cable shall enter the UPS by the bottom right hand side entry:
 - i. Remove RHS UPS cover (remove two top screws make two bottom screws loose),
 - ii. Remove bottom cover plate.

Illustration 7-5:



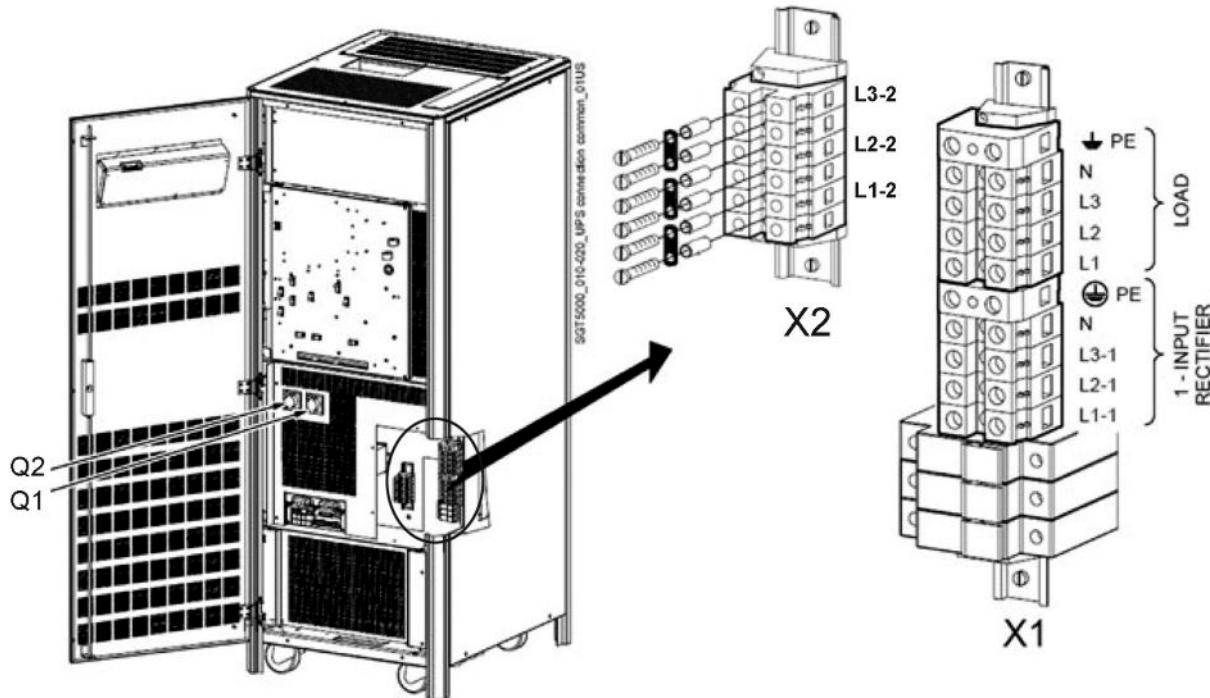
iii. Drill a hole in the bottom cover plate for cable access.



WARNING
RISK OF ELECTRIC SHOCK
HIGH VOLTAGE PRESENT
THE FLUORO UPS REQUIRES COMMON POWER INPUT FOR
RECTIFIER AND BYPASS . JUMPERS SHALL BE KEPT.

e. Check the jumpers linking the terminals L1-2, L2-2 and L3-2, are properly installed.

Illustration 7-6:



f. Connect the Fluoro UPS rectifier input cable from PDB CB5 as follows ([Illustration 7-6](#)):

- Rectifier input phase 1 to X1-L1-1 terminal,
- Rectifier input phase 2 to X1-L2-1 terminal,
- Rectifier input phase 3 to X1-L3-1 terminal,

- Rectifier neutral input to X1-N-rectifier terminal,
 - Rectifier ground to X1-PE-rectifier terminal.
- g. Connect the Fluoro UPS output cable from PDB power block 2 as follows ([Illustration 7-6](#)):
- Output load 1 to X1- L1 terminal,
 - Output load 2 to X1-L2 terminal,
 - Output load 3 to X1-L3 terminal,
 - Ground to X1-PE-Load terminal.
2. Tie the power cables to the UPS and place the protection plates back on the UPS.

1.4 Finalization

No finalization steps.

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