

Interim Facility Site Review Tool

PCP/Clinic Name:				Phone:	Fax/Email:	
Site Address:				Office Contact:		County:
Last Full Scope FSR:	MRR:	FSR Score:	MRR Score:	Health Plan: IEHP R	eviewer:	

Please have the physician or designee complete the form below and return within ten (10) business days. If the completed form is not received within the allotted time frame, an Onsite Interim Review maybe performed. If the answer is "No" to any of the questions, a Corrective Action Plan (CAP) must be submitted to the Health Plan.

Please send completed form and documents to: IEHP QM Coordinator Leonardo-K@iehp.org or (909) 890-5545 (fax).

Check the appropriate Yes/No/NA response below & include any comments. Please note, this form has 2 pages

Physician Coverage is available 24 hours a day, 7 days a week	Compliant	Non- Compliant	N/A	Comments
1. After-hours emergency care instructions/telephone information is made available to patients.	Yes	No		After Hours Access Method: (i.e. phone services/exchange)
CRITICAL ELEMENT	Compliant	Non- Compliant	N/A	Comments
1. Exit doors & aisles are unobstructed and egress (escape) accessible - Accessible pedestrian paths of travel provide a clear circulation path. - Escape routes are maintained free of obstructions or impediments to full instant use of the path of travel in case of fire or other emergency. - Building escape routes provide an accessible, unobstructed path of travel for pedestrians and/or wheelchair users at all times when the site is occupied. - Cords or other items are not placed on or across walkway areas.	Yes	No		
2. Airway Management	Yes	No		Name of person checking supplies:
 Must have a wall oxygen delivery system or portable oxygen tank that is maintained at least ¾ full. Portable oxygen tank must have a flow meter attached. There is a method/system in place for oxygen tank replacement. There are various sizes of oral oropharyngeal airways devices appropriate to patient population available on site. There is a nasal cannula or mask available and various sizes of ambu-bags appropriate to patient population available on site. 				
3. Qualified personnel prepare/administer medication	Yes	No	N/A	Name of MD/NURSE ONLY checking MA
 There must be a licensed physician physically present in the treatment facility during the performance of authorized procedures by the Medical Assistant (MA). There must be a process in place and verbalized by the MA(s), at the time of survey, that the pre-labeled medication container and prepared dose are shown to the licensed person prior to administration. The supervising physician must specifically authorize all medications administered by an MA. 				administered meds:
4. Timely review & follow-up of referral/consultation reports & test results	Yes	No		Please provide referral logs for past 3 months.
 Site staff can demonstrate the office referral process from beginning to end. Referral process must include physician review (e.g. x-ray, labs, specialist notes). A process for follow-up of referral/consultation reports and diagnostic test results is in place. 				Name of person tracking referrals:
 5. Authorized persons dispense medications Drug dispensing is in compliance with all applicable State and Federal laws and regulations. Drugs are dispensed only by a physician, pharmacist or other persons lawfully authorized to dispense medications upon the order of a licensed physician or surgeon. 	Yes	No		Name of MD/NURSE dispensing drugs:
6. Personal protective equipment	Yes	No		
 PPE is available for staff use on site & includes water repelling gloves, water-resistant gowns, face/eye protection (e.g. face shield or goggles), & respiratory infection protection (e.g. mask). 				
7. Needle stick precautions are practiced on site	Yes	No		
 Engineered Sharps Injury Protection (ESIP) devices are used on site Contaminated sharps are discarded immediately. Sharps containers are: 1) located close to the immediate area where sharps are used; 2) inaccessible to unauthorized persons; 3) secured (locked) in patient care areas at all 				
times; and 4) not overfilled past manufacturer's designated fill line or more than ¾ full.				



Interim Facility Site Review Tool

CRITICAL ELEMENT	Compliant	Non- Compliant	N/A	Comments
8. Blood and other infectious materials storage and handling Containers for blood and other potentially infectious materials (OPIM) are closable, leak proof, and labeled and/or color-coded (e.g. red bags). Double bagging is required only if leakage is possible.	Yes	No		
9. Spore testing of autoclave/steam sterilizer • Autoclave spore testing is performed at least monthly. • Written procedures for performing routine spore testing and for handling positive spore test results are available on site to staff. • For positive spore tests, the autoclave is removed from service immediately until inspection is completed and a negative retest occurs. • Procedures include: report problem, repair autoclave, retrieve all instruments sterilized since last negative spore test, re-test autoclave and re-sterilize retrieved instruments.	Yes	No	N/A	<u>Please provide spore test results for last 3 months.</u> Date of last spore test:
Initial Health Assessment (IHA) Reviewed the Initial Health Assessment attached criteria, including how to locate my newly assigned membership on the IEHP website.	Yes	No		

Attestation: I hereby affirm, the information indicated on this form and any documents thereto is true, current, correct and complete to the best of my knowledge and belief, and is furnished in good faith. Lunderstand that material omissions or misrepresentations may result in denial of my application or termination of my

privileges or physician participation agreement	ona onnocon o o mio oproconatorio maj rocat in aonia	or my approaches to termination of my
PCP/Representative Signature & Title:	Date:	
Printed Name & Title:		
MEDICAL GROUP OR HEALTH PLAN USE ONLY		
Interim Review Approved: Yes No	Follow-up Required: Yes No	Date CAP Due:
Nurse Comments:		
Nurse Reviewer Signature:		Date: