preventing tuberculosis (tb) in healthy household contacts of newly diagnosed sputum positive pulmonary TB patients" (1-1060) Project Entitled "A phase III, randomized, double-blind, placebo controlled trial to evaluate the efficacy and safety of vpm1002 and immuvac vaccines in

In lieu of uploaded advertisement for the post of Consultant (Medical Officer), the eligibility criterion is to be as followed:

Post	Consultant (Medical Officer) Essential Qualifica	Post: 1(one) Postgraduate diplotwo years' experience preferance p	_	Desirable Qualific	Desirable Qualific Experience in research /Clinic	Experience in research /Clinica Able to prepare management an supporting them	• Experience in research / Clinica management a supporting then quality systems	• Experience in research /Clinic • Experience in research /Clinic • Able to prepar management a supporting the supporting the • Experience in quality systems • Able to prepar reports and SA	• Experience in research /Clinic • Able to prepare management a supporting the supports and SA • Thorough knoregulatory requirement.
Qualification	Essential Qualification: Post Graduate Degree (MD/MS/DNB) after MBBS with one year experience OR	Postgraduate diploma in medical subjects after MBBS with two years' experience OR MBBS degree with 4 years' experience preferably in clinical research/trial after MBBS		Degree Desirable Qualification and & Experience:	gree sirable Qualification and & Experience: Experience in conducting Vaccine/drug trial/clinicesearch / Clinical Management.	Degree Desirable Qualification and & Experience: Experience in conducting Vaccine/drug trial/clinical research /Clinical Management. Able to prepare safety reports and ensure the timely management and reporting of AEs an SAEs by sites by supporting them	research / Clinical Management. • Experience in conducting Vaccine/drug trial/clinical research / Clinical Management. • Able to prepare safety reports and ensure the timely management and reporting of AEs an SAEs by sites by supporting them • Experience in managing and maintaining databases for quality systems.	egree esirable Qualification and & Experience: Experience in conducting Vaccine/drug trial/clinical research / Clinical Management. Able to prepare safety reports and ensure the timely management and reporting of AEs an SAEs by sites by supporting them Experience in managing and maintaining databases for quality systems. Able to prepare SOPs for trial conduct and write safety reports and SAE narratives.	sirable Qualification and & Experience: Experience in conducting Vaccine/drug trial/clinical management. Able to prepare safety reports and ensure the time management and reporting of AEs an SAEs by sites supporting them Experience in managing and maintaining databases quality systems. Able to prepare SOPs for trial conduct and write safe reports and SAE narratives. Thorough knowledge of GCP, ICH guidelines a regulatory requirements for clinical trial conduct.
Experi	ree Nil	after MBBS with th 4 years'	rial after MBBS	rial after MBBS	mal after MBBS drug trial/clinical	drug trial/clinical ensure the timely SAEs by sites by	rial after MBBS drug trial/clinical ensure the timely SAEs by sites by ling databases for	drug trial/clinical ensure the timely SAEs by sites by ing databases for t and write safety	rnal after MBBS ansure the timely SAEs by sites by ing databases for t and write safety t and write safety I conduct.
Experience									
Salary *per month	70,000/-								
Age Limit	70 years								

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प्रोजेक्ट एंटाइटेल : "एक चरण III, याहच्छिक, डबल-ब्लाइंड, प्लेसबो नियंत्रित परीक्षण vpm1002 की प्रभावकारिता और सुरक्षा का मूल्यांकन करने के लिए और नए निदान किए गए थूक सकारात्मक फुफ्फुसीय टीबी रोगियों के स्वस्थ घरेलू संपर्कों में तपेदिक (टीबी) को रोकने में इम्मुवैक टीके" (I-1060)) 14/10/2021

सलाहकार (चिकित्सा अधिकारी) के पद के लिए अपलोड किए गए विज्ञापन के स्थान पर, पात्रता मानदंड का पालन किया जाना है:

		पदः 1 (एक)	सलाहकार (चिकित्सा अधिकारी)
•टीके/औषधि परीक्षण/नैदानिक अनुसंधान/नैदानिक प्रबंधन के संचालन में अनुभव। •सुरक्षा रिपोर्ट तैयार करने और एई और एसएई को समर्थन देकर समय पर उनका प्रबंधन और रिपोर्टिंग सुनिश्चित करने में सक्षम •गुणवत्ता प्रणालियों के लिए डेटाबेस के प्रबंधन और रिपोर्टिंग खरखाव में अनुभव। •परीक्षण संचालन के लिए एसओपी तैयार करने और सुरक्षा रिपोर्ट और एसएई कथन लिखने में सक्षम। •नैदानिक परीक्षण आचरण के लिए जीसीपी, आईसीएच दिशानिदेशों और नियामक अवश्यकताओं की पूरी जानकारी। •अन्वश संचार कौशल	वांछनीय योग्यता और अनुभवः	एमबीबीएस के बाद पोस्ट ग्रेजुएट डिग्नी (एमडी / एमएस / डीएनबी) एक साल के अनुभव के साथ या एमबीबीएस के बाद मेडिकल विषयों में पोस्टग्रेजुएट डिग्नी के साल के अनुभव के साथ या एमबीबीएस कि जो अनुभव के साथ या एमबीबीएस डिग्नी के साथ 4 साल के अनुभव के साथ एमबीबीएस डिग्नी के बाद क्लिनिकल रिसर्च / ट्रायल में अधिमानतः	योग्यता आवश्यक योग्यताः
		Ž.	अनुभव
		70,000/-	वेतन *प्रति माह
		70 साल	आयु सीमा

