List of Bioequivalence Centres Approved by GCC

No.	Bioequivalence Centres		State
1	Accutest Research Laboratories Pvt Ltd (15 - 18 May 2012), Navi Mumbai – INDIA	2012	Invalid
2	Accutest Research Laboratories (I) Pvt Ltd (Unit-I) (26-27 September 2011), Navi Mumbai -	2011	Invalid
3	Aizant Drug Research Solutions Pvt Ltd (20 - 21 October 2011), Hyderabad - INDIA	2011	Invalid
4	BA Research India Ltd (18-20 October 2010), Ahmedabad - INDIA	2010	Invalid
5	QPS Bioserve India Private Limited - INDIA	2013	Under re-registration
6	Bombay Bioresearch Centre (BBRC) - (18-20 July 2011), Mumbai - INDIA	2011	Invalid
7	Bombay Bioresearch Centre (BBRC) - (18-20 May 2011), Mumbai - INDIA	2011	Invalid
8	Bombay Bioresearch Centre (BBRC) (11-15 October 2010) - INDIA	2010	Invalid
9	Cliantha Research Ltd (19-22 March 2013), Gujarat - INDIA	2013	Under re-registration
10	Cliantha (previously BA Research India Ltd) (19-22 June 2012), Ahmedabad - INDIA	2012	Invalid
11	Lotus Pvt Ltd (25-27 April 2013 - Bangalore; 29-30 April 2013 - Chennai) – INDIA	2013	Invalid
12	Macleods Pharmaceuticals Ltd (8-12 February 2013), Mumbai – INDIA	2013	Invalid
13	Manipal AcuNova KH Clinical Research Centre (2-6 July 2012), Manipal - INDIA	2012	Invalid
14	Matrix Laboratories Ltd (20-21 May 2010), Hyderabad - INDIA	2010	Invalid
15	AnaCipher Clinical Research Organisation- INDIA	2013	Invalid
16	Sitec Labs Pvt Ltd (14-15 February 2012: bioclinical; 16-17 February 2012: bioanalytical),	2012	Under re-registration
17	Veeda Clinical Research Pvt Ltd (14-18 February 2013), Ahmedabad – INDIA	2013	Under re-registration
18	MDS Pharma Services, USA, and Canada	2013	Invalid
19	ACDIMA Center for Bioequivalence & Pharmaceutical Studies, Jordan.	2013	Under re-registration
20	Jordan Center for Pharmaceutical Research (JCPR), Jordan.	2013	Under re-registration
21	Anapharm,Canada.	2013	Invalid
22	Algorithme pharma Inc., Canada.	2013	Invalid
23	International Pharmaceutical Research Center (IPRC), Jordan.	2013	Under re-registration
24	Triumpharma (Clinical Research Centre), Jordan	2014	Under re-registration
25	Pharmaceutical Research Unit (PRU), Jordan	2014	Invalid

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26	Ranbaxy Clinical Pharmacology Unit (CPU) – Noida, India	2015	Invalid
27	Ranbaxy Clinical Pharmacology Unit (CPU) -New Delhi, India	2015	Invalid
28	Ranbaxy Clinical Pharmacology & Pharmacokinetics (CPP) – Gurgaon, India	2015	Invalid
29	Ranbaxy Clinical Pharmacology and Pharmacokinetics (CPP) – Romania	2015	Invalid
30	APL Research Center (Aurbindo Pharma Limited Research Centre -1, India)	2016	Invalid
31	Glenmark Pharmaceuticals Limited, Mahalaxam, Maharashtra, India	2017	Under re-registration
32	3S Pharmacological consultation and Research, Bucharest, Romania.	2017	Invalid
33	In Ventive helath Clinical, Quebec, Canada	2017	Invalid
34	Quinta analytical s.r.o., Czech Republic	2017	Invalid
35	Zi Diligence Bio Center – Egypt	2017	Under re-registration
36	Saudi Ajal – KSA	2017	Invalid
37	Axis Clinicals Limited, Clinical Unit I- India	2017	Invalid
38	Axis Clinicals Limited, Clinical Unit II- India	2017	Invalid
39	RA Chem Pharma Limited, India Clinical Research & Biosciences Division (CRBio)	2018	Valid
40	Pharmasolutions, Egypt	2018	Valid
41	Pharma Medica research Inc – Canada	2018	Valid
42	Bio Clinical Development, Apotex – Canada	2019	Valid
43	Apotex Research Private Limited – India	2019	Valid
44	Anapharm Europe, S.L - Spain	2019	Valid
45	ClinSync Clinical Research PVT - India	2019	Valid
46	International Center for Bioavailability, Pharmaceutical and Clinical research (ICBR)	2022	Valid
47	PT. Clinisindo Laboratories	2022	Valid
48	M/s Analytical Solutions	2022	Valid
49	Quantys Clinical Private Limited	2022	Valid
50	BlueClinical – Investigação e Desenvolvimento em Saúde, Lda.	2023	Valid
51	Enem Nostrum Remedies Pvt. Ltd.	2023	Valid

- Any approved center in last two years can be approved by two of the control authorities: USFDA, Health Canada, TGA, MHRA & EMA) or WHO together with one of the previously mentioned authorities & then the center can be included of the above list.

List of Bioequivalence Centres Approved by GCC

- Centers not included in this list can address the Executive Board and submit a full file about the center for its visit and hence its approval.
- The Executive Board shall apply this mechanism on the new products presented to the Central Registration as of 1st June, 2014.
- The member states shall comply with these criteria at the peripheral level.
- The Executive Board shall post this list on its website, the list will be periodically reviewed and updated.
- Any center has registered in more than 5 years is most Re-register.