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Regulatory Inspections

INDIA

Cliantha Research has been successfully inspected or audited by following regulatory agencies.

AHMEDABAD

April 24 – 28, 2017: Ahmedabad HQ & Sigma by UK MHRA (Clinical & Bio-Analytical)
March 20 – 23, 2017: Ahmedabad HQ by US FDA (Clinical)
March 14 – 17, 2017: Ahmedabad Sigma by US FDA (Clinical)
April 23 – May 3, 2016: Ahmedabad HQ by US FDA (Bio-Analytical)
Jan 18 – 22, 2016: Ahmedabad HQ by US FDA (Clinical)
Mar 23 – Apr 3, 2015: Ahmedabad Sigma by US FDA (Clinical))
May 19 – 23, 2014: Ahmedabad HQ by US FDA (Clinical & Bio-Analytical)
Jan 20 – 24, 2014: Ahmedabad HQ by US FDA (Clinical)
Jul 15 – 18, 2013: Ahmedabad HQ by Thai MoPH (GLP)
Apr 9 – 11, 2013: Ahmedabad HQ and Sigma by Turkey MoH (GLP & GCP)
Mar 19 – 22, 2013: Ahmedabad HQ & Vadodara by WHO (Clinical & Bio-Analytical)
Feb 25 – Mar 1, 2013: Ahmedabad HQ by UK MHRA (Clinical & Bio-Analytical)
Jun 19 – 22, 2012: Ahmedabad HQ and Sigma by WHO (Clinical & Bio-Analytical)
Apr 13 – 15, 2011: Ahmedabad HQ by Spanish Agency (Bio-Analytical)
Oct 25 – 27, 2010: Ahmedabad HQ by US FDA (Clinical)
Oct 18 – 22, 2010: Ahmedabad Sigma by US FDA (Clinical)
Oct 18 – 20, 2010: Ahmedabad HQ by WHO (Clinical & Bio-Analytical)
Jun 17 – 23, 2010: Ahmedabad HQ by US FDA (Bio-Analytical)
Jan 12 – 13, 2009: Ahmedabad HQ by MCC (Clinical & Bio-Analytical)
Sep 20 – 21, 2007: Ahmedabad HQ by US FDA (Clinical & Bio-Analytical)
May 14 – 17, 2007: Ahmedabad HQ by US FDA (Clinical)
Nov 06 – 11, 2006: Ahmedabad HQ by AFSSAPS (ANSM) (Clinical & Bio-Analytical)

VADODARA

Nov 28 – Dec 2, 2016: Vadodara by US FDA (Clinical)
Nov 03 - 05, 2015: Vadodara by AGES (Clinical)
Aug 18 – 22, 2014: Vadodara by US FDA (Clinical)
Jun 24 – 28, 2013: Vadodara by US FDA (Clinical)
Oct 3 – 7, 2011: Vadodara by US FDA (Clinical)
Apr 11 – 12, 2011: Vadodara by Spanish Agency (Clinical)
Feb 9 – 10, 2011: Vadodara by Austrian Agency (Clinical)
Sep 15 – 19, 2008: Vadodara by US FDA (Clinical)

DIAGNOSTIC LAB

Cliantha Sigma: NABL, October 2016
Cliantha Vadodara & Sigma: CAP, September 2015
Cliantha Vadodara & Sigma: CAP, September 2013
Cliantha Sigma: ANVISA, July 2012
Cliantha Vadodara: CAP, August 2011
Cliantha Sigma: CAP, August 2011
Cliantha Vadodara: CAP, September 2010
Cliantha Sigma: CAP, September 2009
Cliantha Vadodara: CAP, September 2008
Cliantha HQ: CAP, September 2007

All the four locations of Cliantha Research Limited - Ahmedabad HQ, Ahmedabad Sigma, Vadodara & Noida (New Delhi) are approved by Drug Controller General of India (DCGI).

NORTH AMERICA

HILL TOP RESEARCH

Hill Top Research, part of Cliantha Research,
has been successfully inspected by following regulatory agencies

2017: Toronto by USFDA (Bio-Analytical) (From May 15 to May 19)
2017: St. Petersburg, FL by USFDA (Clinical) (From May 03 to May 15)
2017: St. Petersburg, FL by USFDA (Clinical) (From Feb 22 to Mar 01)
2016: St. Petersburg, FL by USFDA (Clinical) (From Jan 19 to Feb 19)
2016: St. Petersburg, FL by USFDA (Clinical) (From Jan 11 to Jan 19)
2011: Miamiville, OH by USFDA (Clinical)
2011: St. Petersburg, FL by USFDA (Clinical)
2011: Miamiville, OH by USFDA (Clinical)

INFLAMAX RESEARCH

Inflamax Research, part of Cliantha Research,
has been inspected by both FDA and Health Canada.

2017: Toronto by AEMPS (June)
2016: Toronto by USFDA (November 14 to 17)
2016: Toronto by Health Canada (March 21 to 28)
2016: Toronto by USFDA (March 14 to 17)
2015: Toronto by USFDA (June 15 to 18)
2014: Toronto by USFDA (November 03 to 06)
2014: Toronto by Health Canada (July 21 to 25)
2014: Toronto by Health Canada (March 26)