Regulatory Inspections

INDIA

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

Mar 05 – 09, 2018: Noida by US FDA (Clinical)
Apr 24 – 28, 2017: Ahmedabad HQ & Sigma by UK MHRA (Clinical & Bio-Analytical)
Mar 20 – 23, 2017: Ahmedabad HQ by US FDA (Clinical)
March 14 – 17, 2017: Ahmedabad Sigma by US FDA (Clinical)
Nov 28 – Dec 2, 2016: Vadodara 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)
Nov 03 - 05, 2015: Vadodara by AGES (Clinical)
Mar 23 – Apr 3, 2015: Ahmedabad Sigma by US FDA (Clinical)
Aug 18 – 22, 2014: Vadodara 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)
Jun 24 – 28, 2013: Vadodara by US FDA (Clinical)
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)
Oct 3 – 7, 2011: Vadodara by US FDA (Clinical)
Apr 13 – 15, 2011: Ahmedabad HQ by Spanish Agency (Bio-Analytical)
Apr 11 – 12, 2011: Vadodara by Spanish Agency (Clinical)
Feb 9 – 10, 2011: Vadodara by Austrian Agency (Clinical)
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 15 – 19, 2008: Vadodara by US FDA (Clinical)
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)

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 or audited by following regulatory agencies.

Nov 16 – 20, 2017: Toronto by Spain (AEMPS) (Bio-Analytical)
June 15 2017: Toronto by Health Canada (Bio-Analytical)
May 15 – 19, 2017: Toronto by USFDA (Bio-Analytical)
May 03 – 15, 2017: St. Petersburg, FL by USFDA (Clinical)
Feb 22 – Mar 01, 2017: St. Petersburg, FL by USFDA (Clinical)
Jan 19 – Feb 19, 2016: St. Petersburg, FL by USFDA (Clinical)
Jan 11 – Jan 19, 2016: St. Petersburg, FL by USFDA (Clinical)
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 successfully inspected or audited by following regulatory agencies.

2017: Toronto by AEMPS (June 28 to 30)
2017: Toronto by Health Canada (April 18 to 19)
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)

DIAGNOSTIC LAB

Cliantha Noida: CAP, January 2018
Cliantha Sigma: NABL, October 2017
Cliantha Vadodara & Sigma: CAP, September 2017
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