

**PFIZER INC.**

This document is provided pursuant to Regulation (EC) No 1901/2006 and Communication (2009/C 28/01)

**PROPRIETARY DRUG NAME®/GENERIC DRUG NAME:**

**CHANTIX/VARENICLINE TARTRATE**

**PROTOCOL NO.:**

A3051103

**EudraCT NUMBER:**

**2008-002208-24**

**PROTOCOL TITLE:**

A FOUR WEEK, MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL-GROUP, PHASE 2A STUDY OF VARENICLINE TARTRATE (CP-526,555) IN THE TREATMENT OF POST-HERPETIC NEURALGIA

**Study Centers**

Data not available

**Study Initiation and Final Completion Dates**

Data not available. This study was cancelled prior to enrollment of any study subjects and therefore no data were collected.

**Phase of Development:**

Phase 1

**Study Objectives:**

**1.TO EVALUATE THE ANALGESIC EFFICACY OF VARENICLINE VERSUS PLACEBO IN PATIENTS WITH POSTHERPETIC NEURALGIA (PHN)**

**2.TO EVALUATE THE SAFETY AND TOLERABILITY OF VARENICLINE VERSUS PLACEBO IN PATIENTS WITH POSTHERPETIC NEURALGIA.**

**3.TO CHARACTERIZE THE PHARMACOKINETICS OF VARENICLINE IN PATIENTS WITH POST-HERPETIC NEURALGIA.**

**METHODS**

Data not available

## **RESULTS**

### **Subject Disposition and Demography:**

Data not available

### **Efficacy Results:**

Data not available

### **Safety Results:**

Data not available

## **CONCLUSION:**

Data not available. This study was cancelled prior to enrollment of any study subjects and therefore no data were collected.