Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	**Strength**	**Form**	**Size**	**Date of First Teva Sale***	**Date Discontinued**	**NDC**	**Notes**
Acetaminophen with codeine	300 mg/1, 15 mg/1	TABLET	100	December 2008	Active	00093-0050-01	Previously 00555-0305-02. Changed on 3/22/11
Acetaminophen with codeine	300 mg/1, 30 mg/1	TABLET	100	December 2008	Active	00093-0150-01	Previously 00555-0303-02. Changed on 3/25/10
Acetaminophen with codeine	300 mg/1, 30 mg/1	TABLET	1000	December 2008	Active	00093-0350-10	Previously 00555-0303-05. Changed on 3/25/10
Acetaminophen with codeine	300 mg/1, 60 mg/1	TABLET	100	December 2008	Active	00093-0350-01	Previously 00555-0305-02. Changed on 3/25/10
Acetaminophen with codeine	300 mg/1, 60 mg/1	TABLET	500	December 2008	Active	00093-0350-05	Previously 00555-0304-02. Changed on 11/1/11
Acetaminophen with codeine	300 mg/1, 60 mg/1	TABLET	1000	December 2008	Active	00093-0350-10	Previously 00555-0304-02. Changed on 11/1/11
Actiq	200 ug/1	LOZENGE	30	October 2011	Active	63459-0502-30	
Actiq	400 ug/1	LOZENGE	30	October 2011	Active	63459-0504-30	
Actiq	600 ug/1	LOZENGE	30	October 2011	Active	63459-0506-30	
Actiq	800 ug/1	LOZENGE	30	October 2011	Active	63459-0508-30	
Actiq	1200 ug/1	LOZENGE	30	October 2011	Active	63459-0512-30	
Actiq	1600 ug/1	LOZENGE	30	October 2011	Active	63459-0516-30	
Aspirin (acetylsalicylic acid) with codeine				Exact marketing date unknown, prior to 2006	Discontinued.		
Buprenorphine and Naloxone Sublingual tablets	2 mg/1, .5 mg/1	TABLET	30	December 2014	5/1/2007	00093-5720-56	
Buprenorphine and Naloxone Sublingual tablets	8 mg/1, 2 mg/1	TABLET	30	December 2014	5/1/2007	00093-5721-56	
Buprenorphine and Naloxone Sublingual tablets	8 mg/1, 2 mg/1	TABLET	2 blister pack	August 2016	Active	00228-3155-73	
Buprenorphine and Naloxone Sublingual tablets	2 mg/1, .5 mg/1	TABLET	30	August 2016	Active	00228-3154-73	
Buprenorphine Sublingual tablets	2 mg/1	TABLET	30	May 2010	Teva NDCs temporarily unavailable as of 10/1/2014.	00093-5378-56	
Buprenorphine Sublingual tablets	8 mg/1	TABLET	30	May 2010	Teva NDCs temporarily unavailable as of 10/1/2014.	00093-5379-56	
Buprenorphine Sublingual tablets	2 mg/1	TABLET	30	August 2016	Active	00228-3156-03	
Buprenorphine Sublingual tablets	8 mg/1	TABLET	30	August 2016	Active	00228-3153-03	
Buprenorphine Transdermal System				7/1/2017	Active		
Butalbital, acetaminophen, caffeine, and codeine capsules	50 mg/1, 300 mg/1, 40 mg/1, 30 mg/1	CAPSULE	100	August 2016	Active	00591-2641-01	
Butalbital, acetaminophen, caffeine, and codeine capsules	50 mg/1, 325 mg/1, 40 mg/1, 30 mg/1	CAPSULE	100	August 2016	Active	00591-3220-01	
Butalbital, acetylsalicylic acid, caffeine, and codeine	30 mg/1, 50 mg/1, 40 mg/1, 325 mg/1	CAPSULE	100	August 2016	Active	00591-3546-01	
Butalbital, acetylsalicylic acid, caffeine, and codeine	30 mg/1, 50 mg/1, 40 mg/1, 325 mg/1	CAPSULE	500	August 2016	Active	00591-3546-05	
Fentanyl Transdermal System (Generic Duragesic Patch)	25 ug/h	PATCH, EXTENDED RELEASE	5 pouch	August 2016	Active	00591-3198-72	
Fentanyl Transdermal System (Generic Duragesic Patch)	50 ug/h	PATCH, EXTENDED RELEASE	5 pouch	August 2016	Active	00591-3212-72	
Fentanyl Transdermal System (Generic Duragesic Patch)	75 ug/h	PATCH, EXTENDED RELEASE	5 pouch	August 2016	Active	00591-3213-72	

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PLAINTIFFS TRIAL EXHIBIT
P-18152_00001

Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	Strength	Form	Size	Date of First Teva Sale*	Date Discontinued	NDC	Notes
Fentanyl Transdermal System (Generic Duragesic Patch)	100 ug/h	PATCH, EXTENDED RELEASE	5 pouch	August 2016	Active	00591-3214-72	
Fentora	100 ug/1	TABLET	1 Tablet	10/1/2011	Active	63459-0541-04	
Fentora	100 ug/1	TABLET	28 Tablets	10/1/2011	Active	63459-0541-28	
Fentora	200 ug/1	TABLET	1 Tablet	10/1/2011	Active	63459-0542-04	
Fentora	200 ug/1	TABLET	28 Tables	10/1/2011	Active	63459-0542-28	
Fentora	400 ug/1	TABLET	1 Tablet	10/1/2011	Active	63459-0544-04	
Fentora	400 ug/1	TABLET	28 Tablets	10/1/2011	Active	63459-0544-28	
Fentora	600 ug/1	TABLET	1 Tablet	10/1/2011	Active	63459-0546-04	
Fentora	600 ug/1	TABLET	28 Tablets	10/1/2011	Active	63459-0546-28	
Fentora	800 ug/1	TABLET	1 Tablet	10/1/2011	Active	63459-0548-04	
Fentora	800 ug/1	TABLET	28 Tablets	10/1/2011	Active	63459-0548-28	
Fioricet® with Codeine (butalbital, acetaminophen, caffeine, and codeine phosphate) capsules	50 mg/1, 300 mg/1, 40 mg/1, 30 mg/1	CAPSULE	100	August 2016	Active	52544-0082-01	
Guiatuss AC Syrup, CV (Sugar Free) (OTC)	100 mg/10 mg/5 mL	Syrup	118 mL	8/1/1982	5/1/2007	00182-0017-37	
Guiatuss AC Syrup, CV (Sugar Free) (OTC)	100 mg/10 mg/5 mL	Syrup	473 mL	8/1/1982		00182-0017-40	
Guiatuss DAC Syrup, CV (Sugar Free) (OTC)	100 mg/30 mg/10 mg/5 mL	Syrup	473 mL	4/1/2005	5/1/2007	00182-1378-40	
Hydrocodone Bitartrate and Acetaminophen tablets	5 mg/1, 300 mg/1	TABLET	100	August 2016	Active	00591-2174-01	
Hydrocodone Bitartrate and Acetaminophen tablets	5 mg/1, 300 mg/1	TABLET	500	August 2016	Active	00591-2174-05	
Hydrocodone Bitartrate and Acetaminophen tablets	7.5 mg/1, 300 mg/1	TABLET	100	August 2016	Active	00591-2175-01	
Hydrocodone Bitartrate and Acetaminophen tablets	7.5 mg/1, 300 mg/1	TABLET	500	August 2016	Active	00591-2175-05	
Hydrocodone Bitartrate and Acetaminophen tablets	10 mg/1, 300 mg/1	TABLET	100	August 2016	Active	00591-2176-01	
Hydrocodone Bitartrate and Acetaminophen tablets	10 mg/1, 300 mg/1	TABLET	500	August 2016	Active	00591-2176-05	
Hydrocodone Bitartrate and Acetaminophen tablets	2.5 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00591-2171-01	
Hydrocodone Bitartrate and Acetaminophen tablets	5 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00591-2172-01	
Hydrocodone Bitartrate and Acetaminophen tablets	5 mg/1, 325 mg/1	TABLET	500	August 2016	Active	00591-2172-05	
Hydrocodone Bitartrate and Acetaminophen tablets	7.5 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00591-2605-01	
Hydrocodone Bitartrate and Acetaminophen tablets	7.5 mg/1, 325 mg/1	TABLET	500	August 2016	Active	00591-2605-05	
Hydrocodone Bitartrate and Acetaminophen tablets	10 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00591-2612-01	
Hydrocodone Bitartrate and Acetaminophen tablets	10 mg/1, 325 mg/1	TABLET	500	August 2016	Active	00591-2612-05	
Hydrocodone Bitartrate and Ibuprofen tablets (Generic Vicoprofen)				April 2003	10/2/2014	00093-5161-01	
Hydrocodone Bitartrate and Ibuprofen tablets (Generic Vicoprofen)	7.5 mg/1, 200 mg/1	TABLET, FILM COATED	100	August 2016	Active	62037-0524-01	
Hydrocodone Bitartrate and Ibuprofen tablets (Generic Vicoprofen)	7.5 mg/1, 200 mg/1	TABLET, FILM COATED	500	August 2016	Active	62037-0524-05	
Hydromet	5 mg/5mL, 1.5 mg/5mL	SOLUTION	473 mL	August 2016	Active	0472-1030-16	
Hydromorphone Hydrochlorice Extended-Release tablets	8 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-3629-01	

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Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	Strength	Form	Size	Date of First Teva Sale*	Date Discontinued	NDC	Notes
Hydromorphone Hydrochloride Extended-Release tablets	12 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-3739-01	
Hydromorphone Hydrochloride Extended-Release tablets	16 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-3630-01	
Hydromorphone Hydrochloride Extended-Release tablets	32 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-3631-01	
Hydromorphone Hydrochloride Injection	10 mg/mL	Injection Solution	10 VIAL in 1 CARTON	January 2011	Active	00703-0110-03	
Hydromorphone Hydrochloride Injection	10 mg/mL	Injection Solution	1 vial in 2 carton	January 2011	Active	00703-0018-01	
Hydromorphone Hydrochloride Injection	10 mg/mL	Injection Solution	10 VIAL in 1 CARTON	January 2011	Active	00703-0113-03	
Hydromorphone Hydrochloride tablets				9/1/1991	Actavis discontinued prior to Teva acquisition		
Meperidine hydrochloride tablets	50 mg/1	Tablet	100	February 1997	Active	00555-0381-02	
Meperidine hydrochloride tablets	100 mg/1	TABLET	100	February 1997	Active	00555-0392-02	
Meperidine/Promethazine capsules				10/1/2000	Actavis discontinued prior to Teva acquisition		
Morphine Sulfate Extended-Release capsules	10 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3501-06	
Morphine Sulfate Extended-Release capsules	20 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3502-06	
Morphine Sulfate Extended-Release capsules	30 mg/ 1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3503-06	
Morphine Sulfate Extended-Release capsules	50 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3504-06	
Morphine Sulfate Extended-Release capsules	60 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3505-06	
Morphine Sulfate Extended-Release capsules	80 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3506-06	
Morphine Sulfate Extended-Release capsules	100 mg/1	CAPSULE, EXTENDED RELEASE	60	August 2016	Active	00228-3507-06	

Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	Strength	Form	Size	Date of First Teva Sale*	Date Discontinued	NDC	Notes
Morphine Sulfate Extended-Release capsules	30 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3090-11	
Morphine Sulfate Extended-Release capsules	45 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3116-11	
Morphine Sulfate Extended-Release capsules	60 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3091-11	
Morphine Sulfate Extended-Release capsules	75 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3117-11	
Morphine Sulfate Extended-Release capsules	90 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3092-11	
Morphine Sulfate Extended-Release capsules	120 mg/1	CAPSULE, EXTENDED RELEASE	100	August 2016	Active	00228-3093-11	
Morphine Sulfate Extended-Release tablets	15 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00228-4270-11	
Morphine Sulfate Extended-Release tablets	30 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00228-4271-11	
Morphine Sulfate Extended-Release tablets	60 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00228-4311-11	
Morphine Sulfate Extended-Release tablets	100 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00228-4323-11	
Morphine Sulfate Extended-Release tablets	200 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00228-4347-11	
Nalbuphine hydrochloride injection				6/1/2007	7/1/2010		
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	200 ug/1	LOZENGE	30	12/1/2015	Active	00093-7865-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	400 ug/1	LOZENGE	30	12/1/2015	Active	00093-7866-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	600 ug/1	LOZENGE	30	12/1/2015	Active	00093-7867-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	800 ug/1	LOZENGE	30	12/1/2015	Active	00093-7868-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	1200 ug/1	LOZENGE	30	12/1/2015	Active	00093-7869-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	1600 ug/1	LOZENGE	30	12/1/2015	Active	00093-7870-65	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	800 ug/1	LOZENGE	1 lozenge	12/1/2015	Active	00093-7868-19	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	800 ug/1	LOZENGE	30	August 2016	Active	55253-0073-30	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)				August 2016	Active	55253-0071-30	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)				August 2016	Active	55253-0072-30	

Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	**Strength**	**Form**	**Size**	**Date of First Teva Sale***	**Date Discontinued**	**NDC**	**Notes**
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)	200 ug/1	LOZENGE	30	August 2016	Active	55253-0070-30	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)				August 2016	Active	55253-0074-30	
Oral Transmucosal Fentanyl Citrate (OTFC) Lozenges (Generic Actic)				August 2016	Active	55253-0075-30	
Oxycodone and Acetaminophen tablets (Generic Percocet)	5 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00228-2981-11	
Oxycodone and Acetaminophen tablets (Generic Percocet)	5 mg/1, 325 mg/1	TABLET	500	August 2016	Active	00228-2981-50	
Oxycodone and Acetaminophen tablets (Generic Percocet)	7.5 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00228-2982-11	
Oxycodone and Acetaminophen tablets (Generic Percocet)	10 mg/1, 325 mg/1	TABLET	100	August 2016	Active	00228-2983-11	
Oxycodone Hydrochloride and Aspirin tablets				August 2016	Active	00591-3551-01	
Oxycodone Hydrochloride and Ibuprofen tablets	5 mg/1, 400 mg/1	TABLET, FILM COATED	100	August 2016	Active	00228-4029-11	
Oxycodone Hydrochloride Extended-Release tablets (Generic OxyContin)	4.8355 mg/1, 325 mg/1	TABLET	100	October 2015	Active	00093-5731-01	
Oxycodone Hydrochloride Extended-Release tablets (Generic OxyContin)	10 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	October 2015	Active	00093-5731-01	
Oxycodone Hydrochloride Extended-Release tablets (Generic OxyContin)	40 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-2693-01	
Oxycodone Hydrochloride Extended-Release tablets (Generic OxyContin)	80 mg/1	TABLET, FILM COATED, EXTENDED RELEASE	100	August 2016	Active	00591-2708-01	
Oxycodone Hydrochloride tablets (Generic Roxicodone)	5 mg/1	TABLET	100	August 2016	Active	00228-2876-11	
Oxycodone Hydrochloride tablets (Generic Roxicodone)	15 mg/1	TABLET	100	August 2016	Active	00228-2878-11	
Oxycodone Hydrochloride tablets (Generic Roxicodone)	30 mg/1	TABLET	100	August 2016	Active	00228-2879-11	
Oxymorphone hydrochloride	10 mg/1	TABLET	100	April 2013	Active, temporarily unavailable as of 2/1/2017	00093-5862-01	
Oxymorphone hydrochloride	5 mg/1	TABLET	100	April 2013	Active, temporarily unavailable as of 2/1/2017	00093-5861-01	
Oxymorphone hydrochloride extended release	10 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3228-06	
Oxymorphone hydrochloride extended release	10 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3228-11	
Oxymorphone hydrochloride extended release	15 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3262-06	
Oxymorphone hydrochloride extended release	15 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3262-11	
Oxymorphone hydrochloride extended release	20 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3229-06	

Teva Pharmaceuticals USA List of Opioid Containing Products							
Product Name	**Strength**	**Form**	**Size**	**Date of First Teva Sale***	**Date Discontinued**	**NDC**	**Notes**
Oxymorphone hydrochloride extended release	20 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3229-11	
Oxymorphone hydrochloride extended release	30 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3263-06	
Oxymorphone hydrochloride extended release	30 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3263-11	
Oxymorphone hydrochloride extended release	40 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3230-06	
Oxymorphone hydrochloride extended release	40 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3230-11	
Oxymorphone hydrochloride extended release	5 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3227-06	
Oxymorphone hydrochloride extended release	5 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3227-11	
Oxymorphone hydrochloride extended release	7.5 mg/1	TABLET, EXTENDED RELEASE	60	August 2016	Discontinued 11/2016	00228-3261-06	
Oxymorphone hydrochloride extended release	7.5 mg/1	TABLET, EXTENDED RELEASE	100	August 2016	Discontinued 11/2016	00228-3261-11	
Pentazocine/acetaminophen tablets				7/1/2000	Actavis discontinued prior to Teva acquisition		
Pentazocine/naloxone tablets	50 mg/1, .5 mg/1	TABLET	100	August 2016	Active	00591-0395-01	
Propoxyphene hydrochloride capsules				7/1/1980	Discontinued prior to 2006		
Propoxyphene napsylate and acetaminophen tablets				7/1/1985	Discontinued prior to 2006		
Tramadol hydrochloride tablets	50 mg/1	TABLET, FILM COATED	100	June 2002	Active	00093-0058-01	
Tramadol hydrochloride tablets	50 mg/1	TABLET, FILM COATED	500	June 2002	Active	00093-0058-05	
Tramadol/acetaminophen tablets				January 2006	Discontinued 11/8/2013	00172-6359-00	Marketing Start: 4/15/2005 – this was launched by Ivax prior to Teva's acquisition in 1/2006
Tramadol/acetaminophen tablets				January 2006	Discontinued 11/8/2013	00172-6359-10	Marketing Start: 4/15/2005 – this was launched by Ivax prior to Teva's acquisition in 1/2006
Tramadol/acetaminophen tablets				January 2006	Discontinued 11/8/2013	00172-6359-60	Marketing Start: 4/15/2005 – this was launched by Ivax prior to Teva's acquisition in 1/2006
Tramadol/acetaminophen tablets				January 2006	Discontinued 11/8/2013	00172-6359-70	Marketing Start: 3/13/2006