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Drug Details

| | |
|---|--|
| Drug Name(s) | BUNAVAIL |
| FDA Application No. | (NDA) 205637 |
| Active Ingredient(s) | BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE |
| Company | BIODELIVERY SCI INTL |
| Original Approval or Tentative Approval Date | June 6, 2014 |
| Chemical Type | 3 New dosage form |
| Review Classification | S Standard review drug |

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #205637

Click on a column header to re-sort the table:

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLDTE Code |
|---------------------------|---|------------------------------|-----------------------------------|----------------------------------|----------------------------|
| BUNAVAIL | BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE | EQ 2.1MG BASE; EQ 0.3MG BASE | FILM;BUCCAL | Prescription No | None |
| BUNAVAIL | BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE | EQ 4.2MG BASE; EQ 0.7MG BASE | FILM;BUCCAL | Prescription No | None |
| BUNAVAIL | BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE | EQ 6.3MG BASE; EQ 1MG BASE | FILM;BUCCAL | Prescription Yes | None |

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