

# IT'S ABOUT TIME.

Pr **addyi**<sup>®</sup>  
(flibanserin)  
100mg, 30 tablets



ADDYI is indicated for the treatment of premenopausal and naturally postmenopausal women  $\leq 60$  years of age with acquired, generalized hypoactive sexual desire disorder (HSDD), as characterized by low sexual desire for a minimum of 6 months, which occurs 75-100% of the time, causes marked distress or interpersonal difficulty and is NOT due to:

- A co-existing medical or psychiatric condition,
- Problems within the relationship, or
- The effects of a medication or other drug substance.

## ADDYI should not be prescribed to any patients with a NO response for questions 1-5

- |   |                           |                          |
|---|---------------------------|--------------------------|
| 1. In the past, was your level of sexual desire or interest good and satisfying to you? | <input type="radio"/> Yes | <input type="radio"/> No |
| 2. Has there been a decrease in your level of sexual desire or interest?                | <input type="radio"/> Yes | <input type="radio"/> No |
| 3. Has sexual desire been decreased for over 6 months, 75-100% of the time?             | <input type="radio"/> Yes | <input type="radio"/> No |
| 4. Are you bothered by your decreased level of sexual desire or interest?               | <input type="radio"/> Yes | <input type="radio"/> No |
| 5. Would you like your level of sexual desire or interest to increase?                  | <input type="radio"/> Yes | <input type="radio"/> No |

Clinical judgment should be used if prescribing ADDYI to patients with a YES response to any of the questions below, as a primary diagnosis other than HSDD may be present.

- |  |                           |                          |
|--|---------------------------|--------------------------|
| 1. Are there any factors that may be contributing to your current decrease in sexual desire or interest? |                           |                          |
| A. An operation, depression, injuries, or other medical condition  | <input type="radio"/> Yes | <input type="radio"/> No |
| B. Medications (ex. SSRIs), drugs, or level of alcohol you are currently taking                          | <input type="radio"/> Yes | <input type="radio"/> No |
| C. Recent childbirth, menopausal symptoms  | <input type="radio"/> Yes | <input type="radio"/> No |
| D. Other sexual issues you may be having (pain, decreased arousal or orgasm)                             | <input type="radio"/> Yes | <input type="radio"/> No |
| E. Your partner's sexual problems  | <input type="radio"/> Yes | <input type="radio"/> No |
| F. Dissatisfaction with your relationship or partner   | <input type="radio"/> Yes | <input type="radio"/> No |
| G. Stress or fatigue   | <input type="radio"/> Yes | <input type="radio"/> No |

**ADDYI should not be prescribed to any patients with a YES response to the questions below.**

1. Do you have hepatic (liver) impairment?  Yes  No
2. Are you pregnant or breastfeeding?  Yes  No
3. Are you currently using alcohol and have a resting systolic blood pressure <110 mm Hg or diastolic blood pressure <60 mmHg?  Yes  No
4. Are you taking any moderate or strong CYP3A4 inhibitors? Some examples of moderate or strong CYP3A4 inhibitors include: ritonavir, flucanazole, ciprofloxacin, erythromycin, boceprevir, nefazodone and grapefruit juice.  
Please consult your healthcare professional for additional guidance.  Yes  No
5. Do you have hypersensitivity to flibanserin or other components of ADDYI?  Yes  No
6. Do you take P-glycoprotein (P-gp) substrates (digoxin)?  Yes  No

**Patients should know how to take ADDYI to promote safe use of the medication.**

1. ADDYI is dosed at bedtime because administration during waking hours increases the risks of hypotension, syncope, and CNS depression (such as somnolence and sedation).  Completed
2. The importance of limiting your alcohol intake in combination with ADDYI, as the combination of ADDYI and alcohol may increase the risk of severe low blood pressure.  Completed
3. You should not drive, operate machinery or engage in activities that require clear thinking until 6 hours after taking ADDYI and until you know how it may affect you.  Completed
4. If you miss a dose, you should skip that dose and take your next dose at bedtime of the following day. Increased doses are not associated with greater effect and can increase the risks of side effects.  Completed
5. ADDYI must be used with caution in patients with any pre-existing cardiovascular conditions. Side effects from ADDYI use have included tachycardia, palpitations, hypotension and syncope.  Completed
6. Not all women benefit from treatment with ADDYI. Treatment should be discontinued after 8 weeks if there has been no improvement.  Completed
7. You give consent to treatment with ADDYI.  Completed
8. You have been given a copy of the "Patient Handout".  Completed

**For additional information please consult your healthcare provider.**