

EXELON[®] PATCH (RIVASTIGMINE TRANSDERMAL PATCH) PATIENT DIARY/PATIENT REMINDER CARD (CONSISTING OF INSTRUCTIONS FOR USE AND MEDICATION RECORD SHEETS)

This information is essential to ensure the safe and effective use of Exelon Patch

INSTRUCTIONS FOR USE OF EXELON PATCH (RIVASTIGMINE TRANSDERMAL PATCH) ^{1,2}

- These Instructions for Use and Medication Record Sheets are essential to ensure the safe and effective use of Exelon Patch.
- If you have any questions or require more information, please read the package information leaflet that came with the medicine.
- If you are not sure about anything, please ask your doctor or pharmacist.

Important to remember

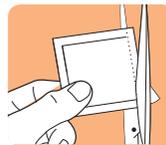
- Take off the previous patch before putting **one** new patch on.
- Only **one** patch per day.
- Do not cut the patch into pieces.
- Press the patch firmly in place for at least 30 seconds using the palm of the hand.

How to apply

1. Carefully remove the existing patch before putting on **one** new patch.



2. Remove the new patch from the sachet.



3. Peel one side of the protective liner off the patch.



4. Stick the patch on the upper or lower back, upper arm, or chest and peel off the protective liner.



5. Press the patch firmly in place for at least 30 seconds.



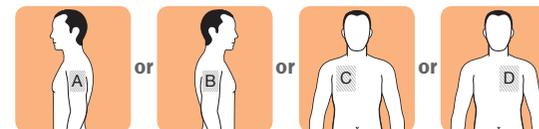
Where to apply

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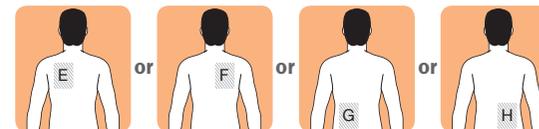
Apply **one** new patch in **one** of the following zones every day.

You can use the same zone (A or B or C or D or E or F or G or H), but do not use the same exact spot within the same zone.

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Back:



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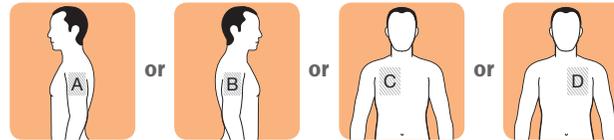
MEDICATION RECORD SHEET FOR EXELON PATCH (RIVASTIGMINE TRANSDERMAL PATCH) ^{1,2}

How to use this record sheet

- Use this record sheet to keep track of when you apply and take off an Exelon Patch.
- Tick the box when you have removed the old patch.
- Fill in the date and day you apply the new patch.
- Fill in the letter of the application zone where you have applied the new patch.
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Application zones

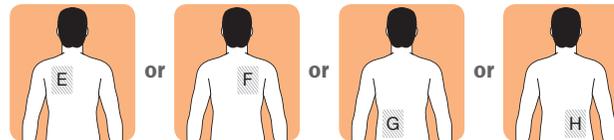
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Apply **one** new patch in **one** of the zones to the left every day.

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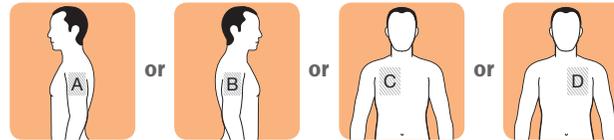
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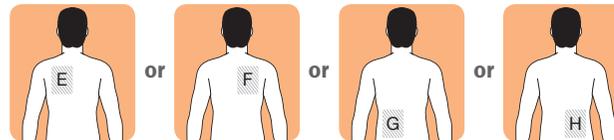
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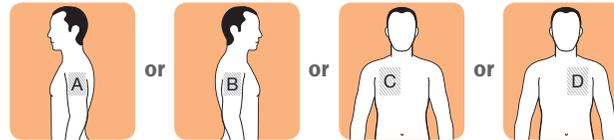
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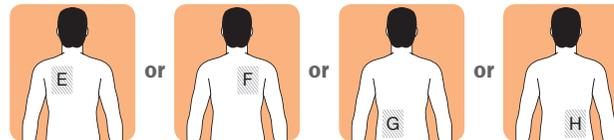
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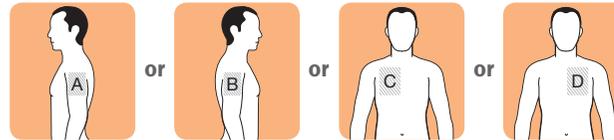
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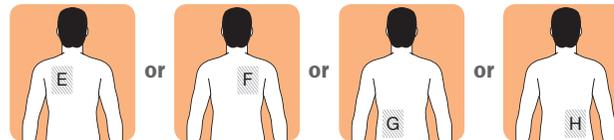
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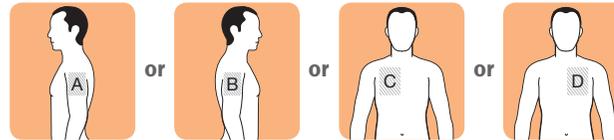
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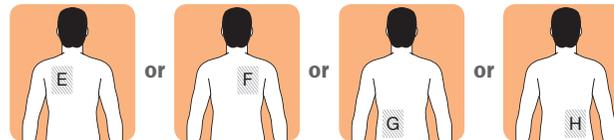
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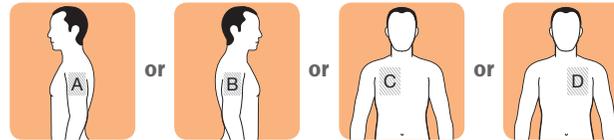
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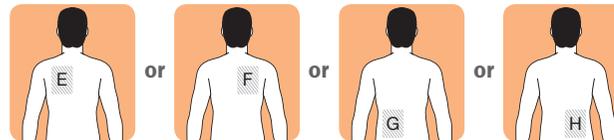
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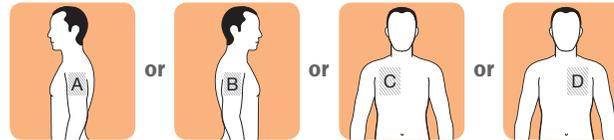
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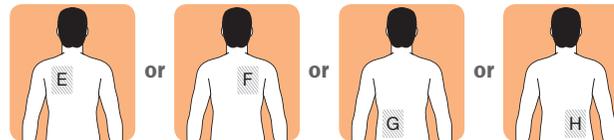
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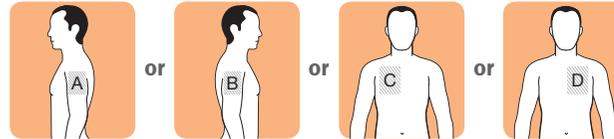
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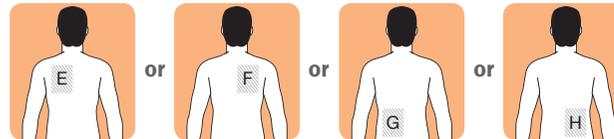
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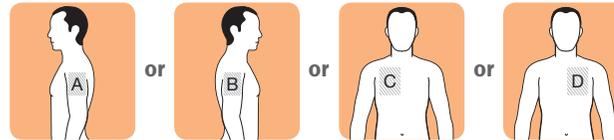
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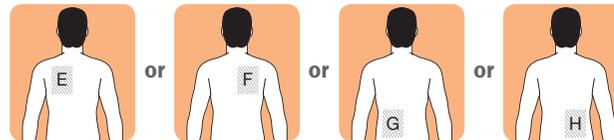
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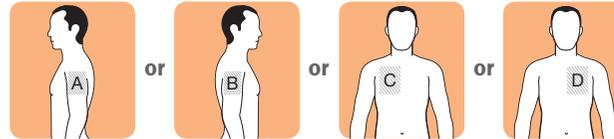
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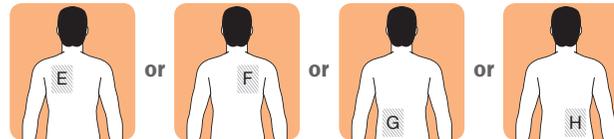
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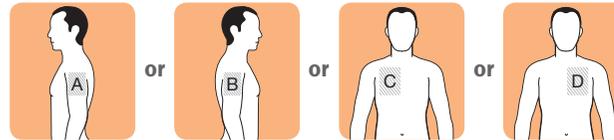
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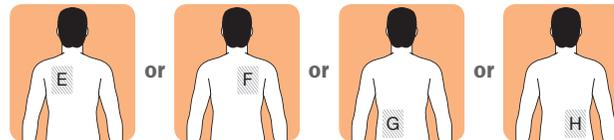
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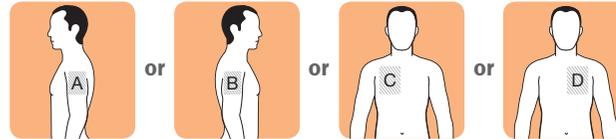
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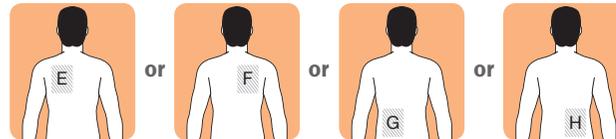
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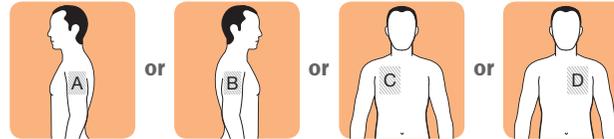
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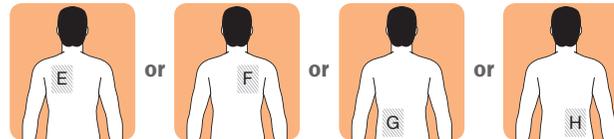
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Suspected adverse reactions and medication errors associated with the use of **Exelon** should be reported to:

Malta Medicines Authority,
Sir Temi Zammit Buildings,
Malta Life Sciences Park,
San Gwann. SGN 3000.

Or at: www.medicinesauthority.gov.mt/adrportal.

Alternatively at: Novartis Pharma Services Inc., Representative Office, Malta by phone on 21222872.

Marketing Authorisation Holder: Novartis Europharm Limited, Frimley Business Park, Camberley GU16 7S4, United Kingdom.

Local Representative: Novartis Pharma Services Inc., Representative Office Malta. Tel No.: +356 21222872.

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EXE PD 04/17 MT

April 2017