

PBR excluded – Funding Application for the use of Botulinum Toxin for Axillary, Palmar or Craniofacial Hyperhidrosis – (form updated February 2016)

Patient NHS No.		Trust:		GP Name:					
Patient Hospital Number:		Consultant Making Request:		GP code / Practice code:					
Patient initials & DoB:		Consultant Contact Details:		GP Post code:					
			Please tick	Please ensure this form is countersigned by Trust Chief Pharmacist (or deputy) before onward transmission to CCG via MECCG.HCD@nhs.net Only fully completed forms will be accepted by Mid-Essex CCG for consideration. If the answer to any of these questions is NO , a full exceptional circumstances form will need to be completed . This may be obtained from the named contact at the relevant PCT/Trust.					
1. Patient has localised hyperhidrosis			<input type="checkbox"/> Yes <input type="checkbox"/> No						
2. Please state which area or areas treatment is being requested for: axillary, palmar or craniofacial			<input type="checkbox"/> Axillary <input type="checkbox"/> Palmar <input type="checkbox"/> Craniofacial						
3. Patient has severe hyperhidrosis and meets the following criteria: <ul style="list-style-type: none"> • Hyperhidrosis Disease Severity Scale (HDSS) > 2 State score: Date:			<input type="checkbox"/> Yes <input type="checkbox"/> No						
4. Patient has tried a strong antiperspirant for at least one month and has not responded to, is intolerant of, or has a contraindication to therapy. Name of antiperspirant: Date started: Date stopped:			<input type="checkbox"/> Yes <input type="checkbox"/> No	<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="height: 40px; vertical-align: top;">Consultant Signature:</td> </tr> <tr> <td style="height: 40px; vertical-align: top;">Approved by Pharmacist/Manager Name:</td> </tr> <tr> <td style="height: 40px; vertical-align: top;">Signature:</td> </tr> <tr> <td style="height: 40px; vertical-align: top;">Date of Application:</td> </tr> </table>		Consultant Signature:	Approved by Pharmacist/Manager Name:	Signature:	Date of Application:
Consultant Signature:									
Approved by Pharmacist/Manager Name:									
Signature:									
Date of Application:									

<p>5. Patient has tried an oral anticholinergic for at least one month and has not responded to, is intolerant of or has a contraindication to therapy.</p> <p>Name of anticholinergic:</p> <p>Date started:</p> <p>Date stopped:</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
<p>6. Patient has tried iontophoresis for at least six months and has not responded to, is intolerant of or has a contraindication to therapy.</p> <p>Date started:</p> <p>Date stopped:</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	

FOR CCG USE ONLY

Funding approvals will be provided for 12 months but injections cannot be given more frequently than once every 6 months i.e. each funding approval will be for 2 injections only, however notification of each injection must be provided in writing.

In all cases repeat injections will not be funded more frequently than every 6 months – please send notification of repeat injections.

Failure to report subsequent injections will result in automatic withdrawal of funding.

Hyperhidrosis Disease Severity Scale (HDSS)

Subjective Score	Clinical interpretation
My sweating is never noticeable and never interferes with my daily activities	1 mild
My sweating is tolerable but sometimes interferes with my daily activities	2 moderate
My sweating is barely tolerable and frequently interferes with my daily activities	3 Severe
My sweating is intolerable and always interferes with my daily activities	4 Severe

From:
Solish N, Benohanian A, Kowalski JW, Canadian Dermatology Study Group on Health-Related Quality of Life in Primary Axillary Hyperhidrosis. Prospective open-label study of botulinum toxin type A in patients with axillary hyperhidrosis: effects on functional impairment and quality of life. *Dermatol Surg* 2005; 31: 405-13.