

AREA PRESCRIBING COMMITTEE – Birmingham, Sandwell, Solihull and environs

Decision Making Support Tool

The following document supports the committee to consider formulary applications against defined criteria.

Formulary application reference:		APCBSSE/0091
Drug name and formulations:		Lisdexamfetamine capsules
Criteria	Example	Committee Consensus
Patient Safety	<i>Potential for abuse, toxicity, significant drug interactions</i>	Theoretically less abusable than other ADHD drugs
Clinical effectiveness	<i>Established licensed product</i>	Equal to or superior to other agents
Strength of evidence		Consistent evidence across all studies
Patient factors	<i>Published patient factors</i>	Once daily dosing schedule
Cost effectiveness or resource impact	£	Cost neutral
Place of therapy relative to available treatments	<i>1/2nd tier</i>	Lisdexamfetamine or methylphenidate 1st line in adults. lisdexamfetamine 2nd line in children over 5 years
National guidance and priorities	<i>NICE, MTRAC</i>	NICE guideline 72: Diagnosis and management of ADHD in children, young people and adults
Local health priorities	<i>CCG views</i>	As per other ADHD agents
Equity of access	<i>Equality assessment</i>	As per other ADHD agents
Stakeholder views	<i>Define wider groups to be engaged</i>	Supportive as per other ADHD agents – subject to commissioning discussions underway
Implementation requirements	<i>Requires, RICAD ESCA etc.</i>	ESCA

Decision Summary

Resubmission is recommended to complete the information to enable a decision:	
Not approved and rationale:	
Formulary status (RAG) and rationale	AMBER with ESCA (to be developed). Please note that commissioning discussion is underway and arrangements currently vary across BSSE area.
Implementation requirements:	ESCA

Implementation monitoring:	