



**Tabla S3. Resumen efectos adversos observados con apremilast en ensayos clínicos y vida real**

<b>ESTEEM-1 y ESTEEM-2<sup>6,9-15</sup> (1)</b>	
<b>EA comunes (≥ 5% de los pacientes) (2)</b>	
Diarrea	18,70%
Náuseas	16,50%
Infección tracto respiratorio superior	19,20%
Nasofaringitis	16,60%
Dolor de cabeza tensional	9,70%
<b>EA graves</b>	
EA cardíacos graves	0,80%
Cáncer	1,90%
Depresión	2,80%
Depresión grave	0,20%
Intención suicida (3)	0,10%
<b>Infecciones</b>	
Tracto urinario	0,20%
Apendicitis	0,30%
Neumonía	0,20%
Otras infecciones	≤ 0,1%
<b>Otros</b>	
Pérdida de peso, media (± SD)	-1,53% (± 6,0%)
<b>Estudio LAPIS-PSO<sup>16</sup>(4)</b>	
<b>EA comunes (≥ 5% de los pacientes)</b>	
Diarrea	8,30%
Náuseas	1,20%
Dolor de cabeza	1,90%
Infección tracto respiratorio superior	0,90%

(1) Seguimiento hasta las 182 semanas. Perfil de seguridad de apremilast en los estudios ESTEEM-1 y ESTEEM-2 comparable al perfil observado en los estudios de fase III LIBERATE y PALACE-1, PALACE-2 y PALACE-37,8.

(2) En general, el porcentaje de pacientes que discontinuaron el tratamiento debido a alguno de estos EA fue muy bajo, ≤1% para cada uno de ellos.

(3) No se registró ningún caso de suicidio consumado.

(4) Parece que la incidencia global de EA con apremilast es menor en la práctica clínica real que la registrada en los ensayos clínicos durante el desarrollo del fármaco.

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